

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
12 December 2002 (12.12.2002)

PCT

(10) International Publication Number
WO 02/098850 A2

(51) International Patent Classification⁷: **C07C 317/46**,
A61K 31/16, A61P 19/02, 33/06, C07C 233/31, 255/29,
255/44, 255/46, 271/12, 271/24, 317/44, C07D 205/04,
207/08, 211/66, 239/08, 263/56, 271/10, C07C 271/16,
C07D 223/08, 277/28, 277/64, 277/82, 295/18, 295/20,
333/24, 307/14, 309/14, 413/12, 413/14, 417/12, 498/04,
C07F 7/18

Gillette, NJ 07933 (US). **THURAIRATNAM, Sukan-
thini** [GB/US]; 6 Smoke Rise Lane, Bedminster, NJ 07921
(US).

(74) Agents: **WANG, George, G.** et al.; Aventis Pharmaceu-
ticals Inc., Route 202-206, P.O. Box 6800, Bridgewater, NJ
08807-0800 (US).

(21) International Application Number: PCT/US02/17411

(22) International Filing Date: 3 June 2002 (03.06.2002)

(25) Filing Language: English

(26) Publication Language: English

(81) Designated States (*national*): AE, AG, AL, AM, AT, AU,
AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU,
CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW,
MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG,
SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ,
VN, YU, ZA, ZM, ZW.

(30) Priority Data:
60/295,301 1 June 2001 (01.06.2001) US

(84) Designated States (*regional*): ARIPO patent (GH, GM,
KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW),
Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM),
European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR,
GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent
(BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR,
NE, SN, TD, TG).

(71) Applicants (*for all designated States except US*): **AXYS
PHARMACEUTICALS, INC.** [US/US]; 180 Kimball
Way, South San Francisco, CA 94080 (US). **AVENTIS
PHARMACEUTICALS INC.** [US/US]; 300 Somerset
Corporate Blvd., Bridgewater, NJ 08807-2854 (US).

(72) Inventors; and

(75) Inventors/Applicants (*for US only*): **GRAUPE, Michael**
[AT/US]; 1131 Banyan Way, Pacifica, CA 94044 (US).
LI, Jiayao [CN/US]; 888 Foster City Blvd., Foster City,
CA 94404 (US). **LINK, John, O.** [US/US]; 12 Prospect
Ave., San Francisco, CA 94110 (US). **ZIPFEL, Sheila**
[US/US]; Ayys Pharmaceuticals Inc., 180 Kimball Way,
South San Francisco, CA 94080 (US). **TIMM, Andreas,**
P. [DE/US]; 97 Bonney Court, Bridgewater, NJ 08807
(US). **ALDOUS, David, J.** [US/US]; 31 Dunbridge Lane,

Declaration under Rule 4.17:

— of inventorship (Rule 4.17(iv)) for US only

Published:

— without international search report and to be republished
upon receipt of that report

*For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
ning of each regular issue of the PCT Gazette.*

WO 02/098850 A2

(54) Title: NOVEL COMPOUNDS AND COMPOSITIONS AS CATHEPSIN INHIBITORS

(57) Abstract: The present invention relates to novel selective cathepsin S inhibitors, the pharmaceutically acceptable salts and N-oxides thereof, their uses as therapeutic agents and the methods of their making.

BEST AVAILABLE COPY

NOVEL COMPOUNDS AND COMPOSITIONS AS CATHEPSIN INHIBITORS

THE INVENTION

5 This application is based on and claims priority from U.S. Provisional Application S.N. 60/295,301 filed on June 1, 2001, incorporated herein by reference.

 This Application relates to compounds and compositions for treating diseases associated with cysteine protease activity, particularly diseases associated with activity of
10 cathepsin S.

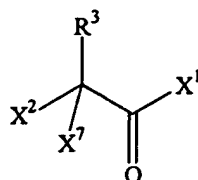
DESCRIPTION OF THE FIELD

 Cysteine proteases represent a class of peptidases characterized by the presence of a
15 cysteine residue in the catalytic site of the enzyme. Cysteine proteases are associated with the normal degradation and processing of proteins. The aberrant activity of cysteine proteases, e.g., as a result of increase expression or enhanced activation, however, may have pathological consequences. In this regard, certain cysteine proteases are associated with a number of disease states, including arthritis, muscular dystrophy, inflammation, tumor invasion,
20 glomerulonephritis, malaria, periodontal disease, metachromatic leukodystrophy and others. An increase in cathepsin S activity contributes to the pathology and/or symptomatology of a number of diseases. Accordingly, molecules that inhibit the activity of cathepsin S protease are useful as therapeutic agents in the treatment of such diseases.

-2-

SUMMARY OF THE INVENTION

This Application relates to compounds of Formula I:



I

in which:

X¹ is -NHC(R¹)(R²)X³ or -NHX⁴;

X² is hydrogen, fluoro, -OH, -OR⁴, -NHR¹⁵ or -NR¹⁷R¹⁸ and X⁷ is hydrogen or X² and X⁷ both represent fluoro;

X³ is cyano, -C(R⁷)(R⁸)R¹⁶, -C(R⁶)(OR⁶)₂, -CH₂C(O)R¹⁶, -CH=CHS(O)₂R⁵, -C(O)CF₂C(O)NR⁵R⁵, -C(O)C(O)NR⁵R⁶, -C(O)C(O)OR⁵, -C(O)CH₂OR⁵, -C(O)CH₂N(R⁶)SO₂R⁵ or -C(O)C(O)R⁵; wherein R⁵ is hydrogen, (C₁-₄)alkyl, (C₃-₁₀)cycloalkyl(C₀-₆)alkyl, hetero(C₃-₁₀)cycloalkyl(C₀-₃)alkyl, (C₆-₁₀)aryl(C₀-₆)alkyl, hetero(C₅-₁₀)aryl(C₀-₆)alkyl, (C₉-₁₀)bicycloaryl(C₀-₆)alkyl or hetero(C₈-₁₀)bicycloaryl(C₀-₆)alkyl; R⁶ is hydrogen, hydroxy or (C₁-₆)alkyl; or where X³ contains an -NR⁵R⁶ group, R⁵ and R⁶ together with the nitrogen atom to which they are both attached, form hetero(C₃-₁₀)cycloalkyl, hetero(C₅-₁₀)aryl or hetero(C₈-₁₀)bicycloaryl; R⁷ is hydrogen or (C₁-₄)alkyl and R⁸ is hydroxy or R⁷ and R⁸ together form oxo; R¹⁶ is hydrogen, -X⁴, -CF₃, -CF₂CF₂R⁹ or -N(R⁶)OR⁶; R⁹ is hydrogen, halo, (C₁-₄)alkyl, (C₅-₁₀)aryl(C₀-₆)alkyl or (C₅-₁₀)heteroaryl(C₀-₆)alkyl, with the proviso that when X³ is cyano, then X² is hydrogen, fluoro, -OH, -OR⁴ or -NR¹⁷R¹⁸ and X⁷ is hydrogen or X² and X⁷ both represent fluoro;

X⁴ comprises a heteromonocyclic ring containing 4 to 7 ring member atoms or a fused heterobicyclic ring system containing 8 to 14 ring member atoms and any carbocyclic ketone, iminoketone or thioketone derivative thereof, with the proviso that when -X⁴ is other than a heteromonocyclic ring containing 5 ring member atoms, wherein no more than two of the ring member atoms comprising the ring are heteroatoms, then X² is fluoro, -OH, -OR⁴, -NHR¹⁵ or -NR¹⁷R¹⁸ and X⁷ is hydrogen or X² and X⁷ both represent fluoro;

wherein within R⁵, X³ or X⁴ any alicyclic or aromatic ring system is unsubstituted or substituted further by 1 to 5 radicals independently selected from (C₁-₆)alkyl, (C₁-₆)alkylidene,

-3-

- cyano, halo, halo-substituted(C₁₋₄)alkyl, nitro, -X⁵NR¹²R¹², -X⁵NR¹²C(O)R¹²,
 -X⁵NR¹²C(O)OR¹², -X⁵NR¹²C(O)NR¹²R¹², -X⁵NR¹²C(NR¹²)NR¹²R¹², -X⁵OR¹², -X⁵SR¹²,
 -X⁵C(O)OR¹², -X⁵C(O)R¹², -X⁵OC(O)R¹², -X⁵C(O)NR¹²R¹², -X⁵S(O)₂NR¹²R¹²,
 -X⁵NR¹²S(O)₂R¹², -X⁵P(O)(OR¹²)OR¹², -X⁵OP(O)(OR¹²)OR¹², -X⁵NR¹²C(O)R¹³, -X⁵S(O)R¹³
 5 and -X⁵S(O)₂R¹³ and/or 1 radical selected from -R¹⁴, -X⁵OR¹⁴, -X⁵SR¹⁴, -X⁵S(O)R¹⁴,
 -X⁵S(O)₂R¹⁴, -X⁵C(O)R¹⁴, -X⁵C(O)OR¹⁴, -X⁵OC(O)R¹⁴, -X⁵NR¹⁴R¹², -X⁵NR¹²C(O)R¹⁴,
 -X⁵NR¹²C(O)OR¹⁴, -X⁵C(O)NR¹²R¹², -X⁵S(O)₂NR¹⁴R¹², -X⁵NR¹²S(O)₂R¹⁴,
 -X⁵NR¹²C(O)NR¹⁴R¹² and -X⁵NR¹²C(NR¹²)NR¹⁴R¹², wherein X⁵ is a bond or (C₁₋₆)alkylene;
 R¹² at each occurrence independently is hydrogen, (C₁₋₆)alkyl or halo-substituted(C₁₋₆)alkyl;
 10 R¹³ is (C₁₋₆)alkyl or halo-substituted(C₁₋₆)alkyl; and R¹⁴ is (C₃₋₁₀)cycloalkyl(C₀₋₆)alkyl,
 hetero(C₃₋₁₀)cycloalkyl(C₀₋₃)alkyl, (C₆₋₁₀)aryl(C₀₋₆)alkyl, hetero(C₅₋₁₀)aryl(C₀₋₆)alkyl,
 (C₉₋₁₀)bicycloaryl(C₀₋₆)alkyl or hetero(C₈₋₁₀)bicycloaryl(C₀₋₆)alkyl;
- R¹ is hydrogen or (C₁₋₆)alkyl and R² is selected from a group consisting of hydrogen,
 cyano, -X⁵NR¹²R¹², -X⁵NR¹²C(O)R¹², -X⁵NR¹²C(O)OR¹², -R¹², -X⁵NR¹²C(O)NR¹²R¹²,
 15 -X⁵NR¹²C(NR¹²)NR¹²R¹², -X⁵OR¹², -X⁵SR¹², -X⁵C(O)OR¹², -X⁵C(O)R¹², -X⁵OC(O)R¹²,
 -X⁵C(O)NR¹²R¹², -X⁵S(O)₂NR¹²R¹², -X⁵NR¹²S(O)₂R¹², -X⁵P(O)(OR¹²)OR¹²,
 -X⁵OP(O)(OR¹²)OR¹², -X⁵NR¹²C(O)R¹³, -X⁵S(O)R¹³, -X⁵S(O)₂R¹³, -R¹⁴, -X⁵OR¹⁴, -X⁵SR¹⁴,
 -X⁵S(O)R¹⁴, -X⁵S(O)₂R¹⁴, -X⁵C(O)R¹⁴, -X⁵C(O)OR¹⁴, -X⁵OC(O)R¹⁴, -X⁵NR¹⁴R¹²,
 -X⁵NR¹²C(O)R¹⁴, -X⁵NR¹²C(O)OR¹⁴, -X⁵C(O)NR¹²R¹², -X⁵S(O)₂NR¹⁴R¹², -X⁵NR¹²S(O)₂R¹⁴,
 20 -X⁵NR¹²C(O)NR¹⁴R¹² and -X⁵NR¹²C(NR¹²)NR¹⁴R¹², wherein X⁵, R¹², R¹³ and R¹⁴ are as
 defined above; or R¹ and R² taken together with the carbon atom to which both R¹ and R² are
 attached form (C₃₋₈)cycloalkylene or (C₃₋₈)heterocycloalkylene; wherein within said R² any
 heteroaryl, aryl, cycloalkyl, heterocycloalkyl, cycloalkylene or heterocycloalkylene is
 unsubstituted or substituted with 1 to 3 radicals independently selected from (C₁₋₆)alkyl,
 25 (C₁₋₆)alkylidene, cyano, halo, halo-substituted(C₁₋₄)alkyl, nitro, -X⁵NR¹²R¹², -X⁵NR¹²C(O)R¹²,
 -X⁵NR¹²C(O)OR¹², -X⁵NR¹²C(O)NR¹²R¹², -X⁵NR¹²C(NR¹²)NR¹²R¹², -X⁵OR¹², -X⁵SR¹²,
 -X⁵C(O)OR¹², -X⁵C(O)R¹², -X⁵OC(O)R¹², -X⁵C(O)NR¹²R¹², -X⁵S(O)₂NR¹²R¹²,
 -X⁵NR¹²S(O)₂R¹², -X⁵P(O)(OR¹²)OR¹², -X⁵OP(O)(OR¹²)OR¹², -X⁵NR¹²C(O)R¹³, -X⁵S(O)R¹³,
 -X⁵S(O)₂R¹³ and -X⁵C(O)R¹³, wherein X⁵, R¹² and R¹³ are as defined above;
- 30 R³ is (C₁₋₆)alkyl or -C(R⁶)(R⁶)X⁶, wherein R⁶ is hydrogen or (C₁₋₆)alkyl and X⁶ is
 selected from -X⁵NR¹²R¹², -X⁵NR¹²C(O)R¹², -X⁵NR¹²C(O)OR¹², -X⁵NR¹²C(O)NR¹²R¹²,
 -X⁵NR¹²C(NR¹²)NR¹²R¹², -X⁵OR¹², -X⁵SR¹², -X⁵C(O)OR¹², -X⁵C(O)R¹², -X⁵OC(O)R¹²,
 -X⁵C(O)NR¹²R¹², -X⁵S(O)₂NR¹²R¹², -X⁵NR¹²S(O)₂R¹², -X⁵P(O)(OR¹²)OR¹²,

-4-

-X⁵OP(O)(OR¹²)OR¹², -X⁵C(O)R¹³, -X⁵NR¹²C(O)R¹³, -X⁵S(O)R¹³, -X⁵S(O)₂R¹³, -R¹⁴,
 -X⁵OR¹⁴, -X⁵SR¹⁴, -X⁵S(O)R¹⁴, -X⁵S(O)₂R¹⁴, -X⁵C(O)R¹⁴, -X⁵C(O)OR¹⁴, -X⁵OC(O)R¹⁴,
 -X⁵NR¹⁴R¹², -X⁵NR¹²C(O)R¹⁴, -X⁵NR¹²C(O)OR¹⁴, -X⁵C(O)NR¹⁴R¹², -X⁵S(O)₂NR¹⁴R¹²,
 -X⁵NR¹²S(O)₂R¹⁴, -X⁵NR¹²C(O)NR¹⁴R¹² and -X⁵NR¹²C(NR¹²)NR¹⁴R¹² wherein X⁵, R¹², R¹³
 5 and R¹⁴ are as defined above;

R⁴ is selected from -X⁸NR¹²R¹², -X⁸NR¹²C(O)R¹², -X⁸NR¹²C(O)OR¹²,
 -X⁸NR¹²C(O)NR¹²R¹², -X⁸NR¹²C(NR¹²)NR¹²R¹², -X⁸OR¹², -X⁸SR¹², -X⁵C(O)OR¹²,
 -X⁵C(O)R¹², -X⁸OC(O)R¹², -X⁵C(O)NR¹²R¹², -X⁸S(O)₂NR¹²R¹², -X⁸NR¹²S(O)₂R¹²,
 -X⁸P(O)(OR¹²)OR¹², -X⁸OP(O)(OR¹²)OR¹², -X⁵C(O)R¹³, -X⁸NR¹²C(O)R¹³, -X⁸S(O)R¹³,
 10 -X⁸S(O)₂R¹³, -R¹⁴, -X⁸OR¹⁴, -X⁸SR¹⁴, -X⁸S(O)R¹⁴, -X⁸S(O)₂R¹⁴, -X⁵C(O)R¹⁴, -X⁵C(O)OR¹⁴,
 -X⁸OC(O)R¹⁴, -X⁸NR¹⁴R¹², -X⁸NR¹²C(O)R¹⁴, -X⁸NR¹²C(O)OR¹⁴, -X⁵C(O)NR¹⁴R¹²,
 -X⁸S(O)₂NR¹⁴R¹², -X⁸NR¹²S(O)₂R¹⁴, -X⁸NR¹²C(O)NR¹⁴R¹² and -X⁸NR¹²C(NR¹²)NR¹⁴R¹²
 wherein X⁸ is (C₁₋₆)alkylene and X⁵, R¹², R¹³ and R¹⁴ are as defined above, with the proviso
 that when X³ is cyano and X² is -OR⁴, where R⁴ is defined as -R¹⁴, then R¹⁴ is

15 (C₃₋₁₀)cycloalkyl(C₁₋₆)alkyl, hetero(C₃₋₁₀)cycloalkyl(C₁₋₃)alkyl, (C₆₋₁₀)aryl(C₁₋₆)alkyl,
 hetero(C₅₋₁₀)aryl(C₁₋₆)alkyl, (C₉₋₁₀)bicycloaryl(C₁₋₆)alkyl or
 hetero(C₈₋₁₀)bicycloaryl(C₁₋₆)alkyl;

R¹⁵ is (C₆₋₁₀)aryl, hetero(C₅₋₁₀)aryl, (C₉₋₁₀)bicycloaryl or hetero(C₈₋₁₀)bicycloaryl;

R¹⁷ is (C₁₋₆)alkyl, (C₃₋₁₀)cycloalkyl(C₀₋₆)alkyl, hetero(C₃₋₁₀)cycloalkyl(C₀₋₃)alkyl,
 20 (C₆₋₁₀)aryl(C₀₋₆)alkyl, hetero(C₅₋₁₀)aryl(C₀₋₆)alkyl, (C₉₋₁₀)bicycloaryl(C₀₋₆)alkyl or
 hetero(C₈₋₁₀)bicycloaryl(C₀₋₆)alkyl, with the proviso that when X³ is cyano, then R¹⁷ is
 (C₁₋₆)alkyl, (C₃₋₁₀)cycloalkyl(C₁₋₆)alkyl, hetero(C₃₋₁₀)cycloalkyl(C₁₋₆)alkyl,
 (C₆₋₁₀)aryl(C₁₋₆)alkyl, hetero(C₅₋₁₀)aryl(C₁₋₆)alkyl, (C₉₋₁₀)bicycloaryl(C₁₋₆)alkyl or
 hetero(C₈₋₁₀)bicycloaryl(C₁₋₆)alkyl;

25 R¹⁸ is hydrogen, (C₁₋₆)alkyl, (C₃₋₁₀)cycloalkyl(C₀₋₆)alkyl,
 hetero(C₃₋₁₀)cycloalkyl(C₀₋₆)alkyl, (C₆₋₁₀)aryl(C₀₋₆)alkyl, hetero(C₅₋₁₀)aryl(C₀₋₆)alkyl,
 (C₉₋₁₀)bicycloaryl(C₀₋₆)alkyl or hetero(C₈₋₁₀)bicycloaryl(C₀₋₆)alkyl, with the proviso that when
 X³ is cyano, then R¹⁸ is (C₁₋₆)alkyl, (C₃₋₁₀)cycloalkyl(C₁₋₆)alkyl,
 hetero(C₃₋₁₀)cycloalkyl(C₁₋₆)alkyl, (C₆₋₁₀)aryl(C₁₋₆)alkyl, hetero(C₅₋₁₀)aryl(C₁₋₆)alkyl,
 30 (C₉₋₁₀)bicycloaryl(C₁₋₆)alkyl or hetero(C₈₋₁₀)bicycloaryl(C₁₋₆)alkyl; and

wherein within R³, R⁴, R¹⁵, R¹⁷ and R¹⁸ any alicyclic or aromatic ring system is
 unsubstituted or substituted further by 1 to 5 radicals independently selected from (C₁₋₆)alkyl,
 (C₁₋₆)alkylidene, cyano, halo, halo-substituted(C₁₋₄)alkyl, nitro, -X⁵NR¹²R¹², -X⁵NR¹²C(O)R¹²,

-5-

$-X^5NR^{12}C(O)OR^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$,
 $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$,
 $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$,
 $-X^5C(O)R^{13}$ and $-X^5S(O)_2R^{13}$ and/or 1 radical selected from $-R^{14}$, $-X^5OR^{14}$, $-X^5SR^{14}$,
5 $-X^5S(O)R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$, $-X^5OC(O)R^{14}$, $-X^5NR^{14}R^{12}$,
 $-X^5NR^{12}C(O)R^{14}$, $-X^5NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{14}R^{12}$, $-X^5S(O)_2NR^{14}R^{12}$, $-X^5NR^{12}S(O)_2R^{14}$,
 $-X^5NR^{12}C(O)NR^{14}R^{12}$ and $-X^5NR^{12}C(NR^{12})NR^{14}R^{12}$; and within R^3 and R^4 any aliphatic
moiety is unsubstituted or substituted further by 1 to 5 radicals independently selected from
cyano, halo, nitro, $-NR^{12}R^{12}$, $-NR^{12}C(O)R^{12}$, $-NR^{12}C(O)OR^{12}$, $-NR^{12}C(O)NR^{12}R^{12}$,
10 $-NR^{12}C(NR^{12})NR^{12}R^{12}$, $-OR^{12}$, $-SR^{12}$, $-C(O)OR^{12}$, $-C(O)R^{12}$, $-OC(O)R^{12}$, $-C(O)NR^{12}R^{12}$,
 $-S(O)_2NR^{12}R^{12}$, $-NR^{12}S(O)_2R^{12}$, $-P(O)(OR^{12})OR^{12}$, $-OP(O)(OR^{12})OR^{12}$, $-NR^{12}C(O)R^{13}$,
 $-S(O)R^{13}$ and $-S(O)_2R^{13}$; wherein X^5 , R^{12} , R^{13} and R^{14} are as described above, with the proviso
that when X^3 is cyano and X^2 is $-OR^4$, where R^4 is defined as $-R^{14}$, or $-NHR^{18}$, then any
aromatic ring system present within R^{14} or R^{18} is not substituted further by halo,
15 (C_{3-10}) cycloalkyl, hetero (C_{3-10}) cycloalkyl, (C_{6-10}) aryl, hetero (C_{5-10}) aryl, (C_{9-10}) bicycloaryl or
hetero (C_{8-10}) bicycloaryl; with the proviso that only one bicyclic ring structure is present
within R^3 , R^4 or R^{15} ; and the *N*-oxide derivatives, prodrug derivatives, protected derivatives,
individual isomers and mixtures of isomers thereof; and the pharmaceutically acceptable salts
and solvates of such compounds and the *N*-oxide derivatives, prodrug derivatives, protected
20 derivatives, individual isomers and mixtures of isomers thereof.

A second aspect of the invention is a pharmaceutical composition which contains a compound of Formula I or their *N*-oxide derivatives, individual isomers or mixture of isomers thereof, or pharmaceutically acceptable salts thereof, in admixture with one or more suitable excipients.

25 A third aspect of the invention is a method for treating a disease in an animal in which inhibition of
cathepsin S can prevent, inhibit or ameliorate the pathology and/or symptomatology of the disease, which
method comprises administering to the animal a therapeutically effective amount of compound of Formula I or a
N-oxide derivative, individual isomer or mixture of isomers thereof; or a pharmaceutically acceptable salt
thereof.

30 A fourth aspect of the invention is the processes for preparing compounds of Formula I and the *N*-oxide
derivatives, prodrug derivatives, protected derivatives, individual isomers and mixtures of isomers thereof; and
the pharmaceutically acceptable salts thereof.

DETAILED DESCRIPTION OF THE INVENTION

35 Definitions:

Unless otherwise stated, the following terms used in the specification and claims are

defined for the purposes of this Application and have the following meanings.

"Alicyclic" means a moiety characterized by arrangement of the carbon atoms in closed non-aromatic ring structures having properties resembling those of aliphatics and may be saturated or partially unsaturated with two or more double or triple bonds.

"Aliphatic" means a moiety characterized by a straight or branched chain arrangement of the constituent carbon atoms and may be saturated or partially unsaturated with two or more double or triple bonds.

"Alkyl" represented by itself means a straight or branched, saturated or unsaturated, aliphatic radical having the number of carbon atoms indicated (e.g., (C₁₋₆)alkyl includes methyl, ethyl, propyl, isopropyl, butyl, *sec*-butyl, isobutyl, *tert*-butyl, vinyl, allyl, 1-propenyl, isopropenyl, 1-butenyl, 2-butenyl, 3-butenyl, 2-methylallyl, ethynyl, 1-propynyl, 2-propynyl, and the like). Alkyl represented along with another radical (e.g., as in arylalkyl) means a straight or branched, saturated or unsaturated aliphatic divalent radical having the number of atoms indicated or when no atoms are indicated means a bond (e.g., (C₆₋₁₀)aryl(C₀₋₃)alkyl includes phenyl, benzyl, phenethyl, 1-phenylethyl 3-phenylpropyl, and the like).

"Alkylene", unless indicated otherwise, means a straight or branched, saturated or unsaturated, aliphatic, divalent radical having the number of carbon atoms indicated (e.g., (C₁₋₆)alkylene includes methylene (-CH₂-), ethylene (-CH₂CH₂-), trimethylene (-CH₂CH₂CH₂-), tetramethylene (-CH₂CH₂CH₂CH₂-), 2-butenylene (-CH₂CH=CHCH₂-), 2-methyltetramethylene (-CH₂CH(CH₃)CH₂CH₂-), pentamethylene (-CH₂CH₂CH₂CH₂CH₂-) and the like).

"Alkylidene" means a straight or branched saturated or unsaturated, aliphatic, divalent radical having the number of carbon atoms indicated (e.g. (C₁₋₆)alkylidene includes methyldiene (=CH₂), ethylidene (=CHCH₃), isopropylidene (=C(CH₃)₂), propylidene (=CHCH₂CH₃), allylidene (=CH'CH=CH₂), and the like).

"Amino" means the radical -NH₂. Unless indicated otherwise, the compounds of the invention containing amino moieties include protected derivatives thereof. Suitable protecting groups for amino moieties include acetyl, *tert*-butoxycarbonyl, benzyloxycarbonyl, and the like.

"Animal" includes humans, non-human mammals (e.g., dogs, cats, rabbits, cattle, horses, sheep, goats, swine, deer, and the like) and non-mammals (e.g., birds, and the like).

"Aromatic" means a moiety wherein the constituent atoms make up an unsaturated ring system, all atoms in the ring system are *sp*² hybridized and the total number of pi electrons is equal to 4n+2.

"Aryl" means a monocyclic or fused bicyclic ring assembly containing the total number of ring carbon atoms indicated, wherein each ring is comprised of 6 ring carbon atoms and is aromatic or when fused with a second ring forms an aromatic ring assembly. For example, optionally substituted (C₆₋₁₀)aryl as used in this Application includes, but is not limited to, biphenyl-2-yl, 2-bromophenyl, 2-bromocarbonylphenyl, 2-bromo-5-fluorophenyl, 4-*tert*-butylphenyl, 4-carbamoylphenyl, 4-carboxy-2-nitrophenyl, 2-chlorophenyl, 4-chlorophenyl, 3-chlorocarbonylphenyl, 4-chlorocarbonylphenyl, 2-chloro-4-fluorophenyl, 2-chloro-6-fluorophenyl, 4-chloro-2-nitrophenyl, 6-chloro-2-nitrophenyl, 2,6-dibromophenyl, 2,3-dichlorophenyl, 2,5-dichlorophenyl, 3,4-dichlorophenyl, 2-difluoromethoxyphenyl, 3,5-dimethylphenyl, 2-ethoxycarbonylphenyl, 2-fluorophenyl, 2-iodophenyl, 4-isopropylphenyl, 2-methoxyphenyl, 4-methoxyphenyl, 2-methylphenyl,

-7-

3-methylphenyl, 4-methylphenyl, 5-methyl-2-nitrophenyl, 4-methylsulfonylphenyl, naphth-2-yl, 2-nitrophenyl, 3-nitrophenyl, 4-nitrophenyl, 2,3,4,5,6-pentafluorophenyl, phenyl, 2-trifluoromethoxyphenyl, 3-trifluoromethoxyphenyl, 4-trifluoromethoxyphenyl, 2-trifluoromethylphenyl, 3-trifluoromethylphenyl, 4-trifluoromethylphenyl, 2-trifluoromethylsulfonylphenyl, 4-trifluoromethylsulfonylphenyl, and the like.

5 Optionally substituted (C_{6-10})aryl as used in this Application includes 3-acetylphenyl, 3-*tert*-butoxycarbonylaminoethylphenyl, biphenyl-4-yl, 3-hydroxyphenyl, 4-hydroxyphenyl, 3-methoxyphenyl, naphth-2-yl, 3-phenoxyphenyl, phenyl, and the like.

"Bicycloaryl" means a bicyclic ring assembly containing the number of ring carbon atoms indicated, wherein the rings are linked by a single bond or fused and at least one of the rings comprising the assembly is aromatic, and any carbocyclic ketone, thio ketone or iminoketone derivative thereof (e.g., (C_{9-10})bicycloaryl includes cyclohexylphenyl, 1,2-dihydronaphthyl, 2,4-dioxo-1,2,3,4-tetrahydronaphthyl, indanyl, indenyl, 1,2,3,4-tetrahydronaphthyl, and the like).

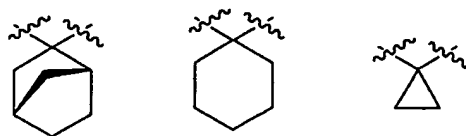
"Carbamoyl" means the radical $-C(O)NH_2$. Unless indicated otherwise, the compounds of the invention containing carbamoyl moieties include protected derivatives thereof. Suitable protecting groups for carbamoyl moieties include acetyl, *tert*-butoxycarbonyl, benzyloxycarbonyl, and the like and both the unprotected and protected derivatives fall within the scope of the invention.

"Carbocyclic ketone derivative" means a derivative containing the moiety $-C(O)-$.

"Carboxy" means the radical $-C(O)OH$. Unless indicated otherwise, the compounds of the invention containing carboxy moieties include protected derivatives thereof. Suitable protecting groups for carboxy moieties include benzyl, *tert*-butyl, and the like.

"Cycloalkyl" means a saturated or partially unsaturated, monocyclic, fused bicyclic or bridged polycyclic ring assembly containing the number of ring carbon atoms indicated, and any carbocyclic ketone, thio ketone or iminoketone derivative thereof (e.g., (C_{3-10})cycloalkyl includes cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cyclohexenyl, 2,5-cyclohexadienyl, bicyclo[2.2.2]octyl, adamantan-1-yl, decahydronaphthyl, oxocyclohexyl, dioxocyclohexyl, thiocyclohexyl, 2-oxobicyclo[2.2.1]hept-1-yl, and the like).

"Cycloalkylene" means a divalent saturated or partially unsaturated, monocyclic ring or bridged polycyclic ring assembly containing the number of ring carbon atoms indicated, and any carbocyclic ketone, thio ketone or iminoketone derivative thereof. For example, the instance wherein R^1 and R^2 together with the carbon atom to which both R^1 and R^2 are attached form (C_{3-8})cycloalkylene" includes, but is not limited to, the following:



35

"Disease" specifically includes any unhealthy condition of an animal or part thereof and includes an unhealthy condition that may be caused by, or incident to, medical or

-8-

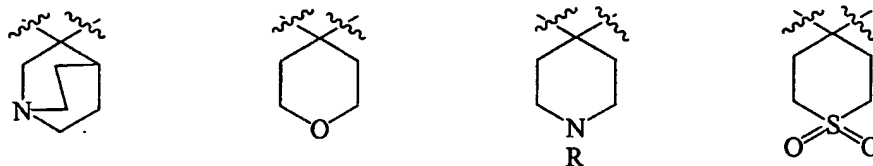
veterinary therapy applied to that animal, i.e., the "side effects" of such therapy.

"Halo" means fluoro, chloro, bromo or iodo.

"Halo-substituted alkyl", as an isolated group or part of a larger group, means "alkyl" substituted by one or more "halo" atoms, as such terms are defined in this Application. Halo-substituted alkyl includes haloalkyl, dihaloalkyl, trihaloalkyl, perhaloalkyl and the like (e.g. halo-substituted (C₁₋₃)alkyl includes chloromethyl, dichloromethyl, difluoromethyl, trifluoromethyl, 2,2,2-trifluoroethyl, perfluoroethyl, 2,2,2-trifluoro-1,1-dichloroethyl, and the like).

"Heteroatom moiety" includes -N=, -NR-, -O-, -S- or -S(O)₂-, wherein R is hydrogen, (C₁₋₆)alkyl or a protecting group.

"Heterocycloalkylene" means cycloalkylene, as defined in this Application, provided that one or more of the ring member carbon atoms indicated, is replaced by heteroatom moiety selected from -N=, -NR-, -O-, -S- or -S(O)₂-, wherein R is hydrogen or (C₁₋₆)alkyl. For example, the instance wherein R¹ and R² together with the carbon atom to which both R¹ and R² are attached form hetero(C₃₋₈)cycloalkyl" includes, but is not limited to, the following:



in which R is hydrogen, (C₁₋₆)alkyl, or a protecting group.

"Heteroaryl" means aryl, as defined in this Application, provided that one or more of the ring carbon atoms indicated are replaced by a heteroatom moiety selected from -N=, -NR-, -O- or -S-, wherein R is hydrogen, (C₁₋₆)alkyl, a protecting group or represents the free valence which serves as the point of attachment to a ring nitrogen, and each ring is comprised of 5 or 6 ring atoms. For example, optionally substituted hetero(C₅₋₁₀)aryl as used in this Application includes, but is not limited to, 4-amino-2-hydroxypyrimidin-5-yl, benzothiazol-2-yl, 1H-benzimidazol-2-yl, 2-bromopyrid-5-yl, 5-bromopyrid-2-yl, 4-carbamoylthiazol-2-yl, 3-carboxypyrid-4-yl, 5-carboxy-2,6-dimethylpyrid-3-yl, 3,5-dimethylisoxazol-4-yl, 5-ethoxy-2,6-dimethylpyrid-3-yl, 5-fluoro-6-hydroxypyrimidin-4-yl, fur-2-yl, fur-3-yl, 5-hydroxy-4,6-dimethylpyrid-3-yl, 8-hydroxy-5,7-dimethylquinolin-2-yl, 5-hydroxymethylisoxazol-3-yl, 3-hydroxy-6-methylpyrid-2-yl, 3-hydroxypyrid-2-yl, 1H-imidazol-2-yl, 1H-imidazol-4-yl, 1H-indol-3-yl, isothiazol-4-yl, isoxazol-4-yl, 2-methylfur-3-yl, 5-methylfur-2-yl, 1-methyl-1H-imidazol-2-yl, 5-methyl-3H-imidazol-4-yl, 5-methylisoxazol-3-yl, 5-methyl-2H-pyrazol-3-yl, 3-methylpyrid-2-yl, 4-methylpyrid-2-yl, 5-methylpyrid-2-yl, 6-methylpyrid-2-yl, 2-methylpyrid-3-yl, 2-methylthiazol-4-yl, 5-nitropyrid-2-yl, 2H-pyrazol-3-yl, 3H-pyrazol-4-yl, pyridazin-3-yl, pyrid-2-yl, pyrid-3-yl, pyrid-4-yl, 5-pyrid-3-yl-2H-[1,2,4]triazol-3-yl, pyrimidin-4-yl, pyrimidin-5-yl, 1H-pyrrol-3-yl, quinolin-2-yl, 1H-tetrazol-5-yl, thiazol-2-yl, thiazol-5-yl, thien-2-yl, thien-3-yl, 2H-[1,2,4]triazol-3-yl, 3H-[1,2,3]triazol-4-yl, 5-trifluoromethylpyrid-2-yl, and the like. Suitable protecting groups include *tert*-butoxycarbonyl, benzyloxycarbonyl, benzyl, 4-methoxybenzyl, 2-nitrobenzyl, and the like.

Optionally substituted hetero(C₅₋₁₀)aryl as used in this Application to define R⁴ includes benzofur-2-yl, fur-2-yl, fur-3-yl, pyrid-3-yl, pyrid-4-yl, quinol-2-yl, quinol-3-yl, thien-2-yl, thien-3-yl, and the like.

"Heterobicycloaryl" means bicycloaryl, as defined in this Application, provided that one or more of the ring carbon atoms indicated are replaced by a heteroatom moiety selected from -N=, -NR-, -O- or -S-, wherein R is hydrogen, (C₁₋₆)alkyl, a protecting group or represents the free valence which serves as the point of attachment to a ring nitrogen, and any carbocyclic ketone, thioketone or iminoketone derivative thereof. For example, optionally substituted hetero(C₈₋₁₀)bicycloaryl as used in this Application includes, but is not limited to, 2-amino-4-oxo-3,4-dihydropteridin-6-yl, and the like. In general, the term heterobicycloaryl as used in this Application includes, for example, benzo[1,3]dioxol-5-yl, 3,4-dihydro-2*H*-[1,8]naphthyridinyl, 3,4-dihydro-2*H*-quinolinyl, 2,4-dioxo-3,4-dihydro-2*H*-quinazolinyl, 1,2,3,4,5,6-hexahydro[2,2']bipyridinyl, 3-oxo-2,3-dihydrobenzo[1,4]oxazinyl, 5,6,7,8-tetrahydroquinolinyl, and the like.

"Heterocycloalkyl" means cycloalkyl, as defined in this Application, provided that one or more of the ring carbon atoms indicated are replaced by a heteroatom moiety selected from -N=, -NR-, -O- or -S-, wherein R is hydrogen, (C₁₋₆)alkyl, a protecting group or represents the free valence which serves as the point of attachment to a ring nitrogen, and any carbocyclic ketone, thioketone or iminoketone derivative thereof (e.g., the term hetero(C₅₋₁₀)cycloalkyl includes imidazolidinyl, morpholinyl, piperazinyl, piperidyl, pyrrolidinyl, pyrrolinyl, quinuclidinyl, and the like). Suitable protecting groups include *tert*-butoxycarbonyl, benzyloxycarbonyl, benzyl, 4-methoxybenzyl, 2-nitrobenzyl, and the like. Both the unprotected and protected derivatives fall within the scope of the invention.

"Heteromonocyclic ring" means a saturated or partially unsaturated, monocyclic ring assembly containing the number of ring carbon atoms indicated, as defined in this Application, provided that one or more of the ring carbon atoms indicated are replaced by one or more heteroatoms selected from -N=, -NY³-, -O- or -S-, wherein Y³ is hydrogen, alkyl, aryl, arylalkyl, -C(=O)-R¹⁴, -C(=O)-OR¹⁴ or -SO₂R¹⁴.

"Heterobicyclic ring" means a saturated or partially unsaturated fused bicyclic or bridged polycyclic ring assembly containing the number of ring carbon atoms indicated, as defined in this Application, provided that one or more of the ring carbon atoms indicated are replaced by one or more heteroatoms selected from -N=, -NY³-, -O- or -S-, wherein Y³ is hydrogen, alkyl, aryl, arylalkyl, -C(=O)-R¹⁴, -C(=O)-OR¹⁴ or -SO₂R¹⁴.

"Hydroxy" means the radical -OH. Unless indicated otherwise, the compounds of the invention containing hydroxy radicals include protected derivatives thereof. Suitable protecting groups for hydroxy moieties include benzyl and the like.

"Iminoketone derivative" means a derivative containing the moiety -C(NR)-, wherein R is hydrogen or (C₁₋₆)alkyl.

"Isomers" mean compounds of Formula I having identical molecular formulae but differ in the nature or sequence of bonding of their atoms or in the arrangement of their atoms in space. Isomers that differ in the arrangement of their atoms in space are termed "stereoisomers". Stereoisomers that are not mirror images of one another are termed "diastereomers" and stereoisomers that are nonsuperimposable mirror images are termed "enantiomers" or sometimes "optical isomers". A carbon atom bonded to four nonidentical substituents is termed a "chiral center". A compound with one chiral center has two enantiomeric forms of opposite chirality is termed a "racemic mixture". A compound that has more than one chiral center has 2^{*n*-1} enantiomeric pairs, where *n* is the number of chiral centers. Compounds with more than one chiral center may exist as either an individual

diastereomers or as a mixture of diastereomers, termed a "diastereomeric mixture". When one chiral center is present a stereoisomer may be characterized by the absolute configuration of that chiral center. Absolute configuration refers to the arrangement in space of the substituents attached to the chiral center. Enantiomers are characterized by the absolute configuration of their chiral centers and described by the *R*- and *S*-sequencing rules of Cahn, Ingold and Prelog. Conventions for stereochemical nomenclature, methods for the determination of stereochemistry and the separation of stereoisomers are well known in the art (e.g., see "Advanced Organic Chemistry", 4th edition, March, Jerry, John Wiley & Sons, New York, 1992). It is understood that the names and illustration used in this Application to describe compounds of Formula I are meant to be encompassed all possible stereoisomers. Thus, for example, the name *N*-[1-(1-benzothiazol-2-yl-methanoyl)-propyl]-2-hydroxy-3-phenylmethanesulfonyl-propionamide is meant to include (S)-*N*-[1-(1-benzothiazol-2-yl-methanoyl)-propyl]-2-hydroxy-3-phenylmethanesulfonyl-propionamide, (R)-*N*-[1-(1-benzothiazol-2-yl-methanoyl)-propyl]-2-hydroxy-3-phenylmethanesulfonyl-propionamide, (R)-*N*-[(S)-1-(1-benzothiazol-2-yl-methanoyl)-propyl]-2-hydroxy-3-phenylmethanesulfonyl-propionamide, (S)-*N*-[(R)-1-(1-benzothiazol-2-yl-methanoyl)-propyl]-2-hydroxy-3-phenylmethanesulfonyl-propionamide, (R)-*N*-[(R)-1-(1-benzothiazol-2-yl-methanoyl)-propyl]-2-hydroxy-3-phenylmethanesulfonyl-propionamide, *N*-[(S)-1-(1-benzothiazol-2-yl-methanoyl)-propyl]-2-hydroxy-3-phenylmethanesulfonyl-propionamide, *N*-[(R)-1-(1-benzothiazol-2-yl-methanoyl)-propyl]-2-hydroxy-3-phenylmethanesulfonyl-propionamide, (S)-*N*-[(S)-1-(1-benzothiazol-2-yl-methanoyl)-propyl]-2-hydroxy-3-phenylmethanesulfonyl-propionamide and any mixture, racemic or otherwise, thereof.

"Ketone derivative" means a derivative containing the moiety -C(O)-. For example, in this Application X³ can be 2-acetoxy-azetidin-3-yl. The "carbocyclic ketone derivative" of this example of X³ would be 2-acetoxy-4-oxo-azetidin-3-yl (see Table 3, C32).

"Nitro" means the radical -NO₂.

"Optional" or "optionally" means that the subsequently described event or circumstance may or may not occur, and that the description includes instances where the event or circumstance occurs and instances in which it does not. For example, the phrase "wherein within R³ and R⁴ any alicyclic or aromatic ring system may be substituted further by 1-5 radicals..." means that R³ and R⁴ may or may not be substituted in order to fall within the scope of the invention.

"Oxoalkyl" means alkyl, as defined above, wherein one of the number of carbon atoms indicated is replaced by an oxygen group (-O-), e.g., oxo(C₂₋₆)alkyl includes methoxymethyl, etc.

"N-oxide derivatives" means derivatives of compounds of Formula I in which nitrogens are in an oxidized state (i.e., O-N) and which possess the desired pharmacological activity.

"Pathology" of a disease means the essential nature, causes and development of the disease as well as the structural and functional changes that result from the disease processes.

"Pharmaceutically acceptable" means that which is useful in preparing a pharmaceutical composition that is generally safe, non-toxic and neither biologically nor otherwise undesirable and includes that which is acceptable for veterinary use as well as human pharmaceutical use.

"Pharmaceutically acceptable salts" means salts of compounds of Formula I which are pharmaceutically acceptable, as defined above, and which possess the desired pharmacological activity. Such salts include acid addition salts formed with inorganic acids such as hydrochloric acid, hydrobromic acid, sulfuric acid, nitric acid, phosphoric acid, and the like; or with organic acids such as acetic acid, propionic acid, hexanoic acid, heptanoic acid, cyclopentanepropionic acid, glycolic acid, pyruvic acid, lactic acid, malonic acid, succinic acid, malic acid, maleic acid, fumaric acid, tartaric acid, citric acid, benzoic acid, *o*-(4-hydroxybenzoyl)benzoic acid, cinnamic acid, maleic acid, methanesulfonic acid, ethanesulfonic acid, 1,2-ethanedisulfonic acid, 2-hydroxyethanesulfonic acid, benzenesulfonic acid, *p*-chlorobenzenesulfonic acid, 2-naphthalenesulfonic acid, *p*-toluenesulfonic acid, camphorsulfonic acid, 4-methylbicyclo[2.2.2]oct-2-ene-1-carboxylic acid, glucoheptonic acid, 4,4'-methylenebis(3-hydroxy-2-ene-1-carboxylic acid), 3-phenylpropionic acid, trimethylacetic acid, tertiary butylacetic acid, lauryl sulfuric acid, gluconic acid, glutamic acid, hydroxynaphthoic acid, salicylic acid, stearic acid, muconic acid and the like.

Pharmaceutically acceptable salts also include base addition salts which may be formed when acidic protons present are capable of reacting with inorganic or organic bases. Acceptable inorganic bases include sodium hydroxide, sodium carbonate, potassium hydroxide, aluminum hydroxide and calcium hydroxide. Acceptable organic bases include ethanolamine, diethanolamine, triethanolamine, tromethamine, *N*-methylglucamine and the like.

"Prodrug" means a compound which is convertible in vivo by metabolic means (e.g. by hydrolysis) to a compound of Formula I. For example an ester of a compound of Formula I containing a hydroxy group may be convertible by hydrolysis in vivo to the parent molecule. Alternatively an ester of a compound of Formula I containing a carboxy group may be convertible by hydrolysis in vivo to the parent molecule. Suitable esters of compounds of Formula I containing a hydroxy group, are for example acetates, citrates, lactates, tartrates, malonates, oxalates, salicylates, propionates, succinates, fumarates, maleates, methylenebis-*b*-hydroxynaphthoates, gentisates, isethionates, di-*p*-toluoyltartrates, methanesulphonates, ethanesulphonates, benzenesulphonates, *p*-toluenesulphonates, cyclohexylsulphamates and quinate. Suitable esters of compounds of Formula I containing a carboxy group, are for example those described by F.J. Leinweber, *Drug Metab. Res.*, 1987, 18, page 379. An especially useful class of esters of compounds of Formula I containing a hydroxy group, may be formed from acid moieties selected from those described by Bundgaard et al., *J. Med. Chem.*, 1989, 32, page 2503-2507, and include substituted (aminomethyl)-benzoates, for example, dialkylamino-methylbenzoates in which the two alkyl groups may be joined together and/or interrupted by an oxygen atom or by an optionally substituted nitrogen atom, e.g. an alkylated nitrogen atom, more especially (morpholino-methyl)benzoates, e.g. 3- or 4-(morpholinomethyl)-benzoates, and (4-alkylpiperazin-1-yl)benzoates, e.g. 3- or 4-(4-alkylpiperazin-1-yl)benzoates.

"Protected derivatives" means derivatives of compounds of Formula I in which a reactive site or sites are blocked with protecting groups. Protected derivatives of compounds of Formula I are useful in the preparation of compounds of Formula I or in themselves may be active cathepsin S inhibitors. A comprehensive list of suitable protecting groups can be found in T.W. Greene, *Protecting Groups in Organic Synthesis*, 3rd edition, John Wiley & Sons, Inc. 1999.

"Therapeutically effective amount" means that amount which, when administered to an animal for treating a disease, is sufficient to effect such treatment for the disease.

-12-

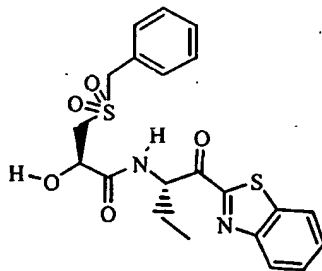
"Thioketone derivative" means a derivative containing the moiety -C(S)-.

"Treatment" or "treating" means any administration of a compound of the present invention and includes:

- (1) preventing the disease from occurring in an animal which may be predisposed to the disease but does not yet experience or display the pathology or symptomatology of the disease,
- (2) inhibiting the disease in an animal that is experiencing or displaying the pathology or symptomatology of the diseased (i.e., arresting further development of the pathology and/or symptomatology), or
- (3) ameliorating the disease in an animal that is experiencing or displaying the pathology or symptomatology of the diseased (i.e., reversing the pathology and/or symptomatology).

Nomenclature:

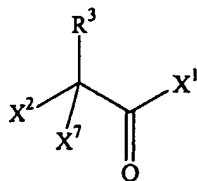
The compounds of Formula I and the intermediates and starting materials used in their preparation are named in accordance with IUPAC rules of nomenclature in which the characteristic groups have decreasing priority for citation as the principle group as follows: acids, esters, amides, etc. Alternatively, the compounds are named by AutoNom 4.0 (Beilstein Information Systems, Inc.). For example, a compound of Formula I wherein X^2 is hydroxy, R^3 is phenylmethanesulfonylmethyl and X^1 is $-NHC(R^1)(R^2)X^3$ (in which R^1 is hydrogen, R^2 is ethyl and X^3 is 1-benzothiazol-2-yl-methanoyl); that is, a compound having the following structure:



is named (R)-N-[(S)-1-(1-benzothiazol-2-yl-methanoyl)-propyl]-2-hydroxy-3-phenylmethanesulfonyl-propionamide;

Presently Preferred Embodiments:

While the scope of the invention is set forth in the Summary of the Invention, certain aspects of the invention are preferred. For example, preferred is a compound of Formula I:



I

in which:

X^1 is $-NHC(R^1)(R^2)X^3$ or $-NHCH(R^{19})C(O)R^{20}$;

-13-

X^2 is hydrogen, fluoro, -OH, -OR⁴, -NHR¹⁵ or -NR¹⁷R¹⁸ and X^7 is hydrogen or X^2 and X^7 both represent fluoro;

X^3 is cyano, -C(R⁷)(R⁸)R¹⁶, -C(R⁶)(OR⁶)₂, -CH₂C(O)R¹⁶, -CH=CHS(O)₂R⁵,
 -C(O)CF₂C(O)NR⁵R⁵, -C(O)C(O)NR⁵R⁶, -C(O)C(O)OR⁵, -C(O)CH₂OR⁵,
 5 -C(O)CH₂N(R⁶)SO₂R⁵ or -C(O)C(O)R⁵; wherein R⁵ is hydrogen, (C₁₋₄)alkyl,
 (C₃₋₁₀)cycloalkyl(C₀₋₆)alkyl, hetero(C₃₋₁₀)cycloalkyl(C₀₋₃)alkyl, (C₆₋₁₀)aryl(C₀₋₆)alkyl,
 hetero(C₅₋₁₀)aryl(C₀₋₆)alkyl, (C₉₋₁₀)bicycloaryl(C₀₋₆)alkyl or
 hetero(C₈₋₁₀)bicycloaryl(C₀₋₆)alkyl; R⁶ is hydrogen, hydroxy or (C₁₋₆)alkyl; or where X^3
 contains an -NR⁵R⁶ group, R⁵ and R⁶ together with the nitrogen atom to which they are both
 10 attached, form hetero(C₃₋₁₀)cycloalkyl, hetero(C₅₋₁₀)aryl or hetero(C₈₋₁₀)bicycloaryl; R⁷ is
 hydrogen or (C₁₋₄)alkyl and R⁸ is hydroxy or R⁷ and R⁸ together form oxo; R¹⁶ is hydrogen, -
 X^4 , -CF₃, -CF₂CF₂R⁹ or -N(R⁶)OR⁶; R⁹ is hydrogen, halo, (C₁₋₄)alkyl, (C₅₋₁₀)aryl(C₀₋₆)alkyl or
 (C₅₋₁₀)heteroaryl(C₀₋₆)alkyl, with the proviso that when X^3 is cyano, then X^2 is hydrogen,
 fluoro, -OH, -OR⁴ or -NR¹⁷R¹⁸ and X^7 is hydrogen or X^2 and X^7 both represent fluoro;

15 X^4 comprises a heteromonocyclic ring containing 4 to 7 ring member atoms or a fused
 heterobicyclic ring system containing 8 to 14 ring member atoms and any carbocyclic ketone,
 iminoketone or thioketone derivative thereof, with the proviso that when - X^4 is other than a
 heteromonocyclic ring containing 5 ring member atoms, wherein no more than two of the ring
 member atoms comprising the ring are heteroatoms, then X^2 is fluoro, -OH, -OR⁴, -NHR¹⁵ or
 20 -NR¹⁷R¹⁸ and X^7 is hydrogen or X^2 and X^7 both represent fluoro;

wherein within R⁵, X^3 or X^4 any alicyclic or aromatic ring system is unsubstituted or
 substituted further by 1 to 5 radicals independently selected from (C₁₋₆)alkyl, (C₁₋₆)alkylidene,
 cyano, halo, halo-substituted(C₁₋₄)alkyl, nitro, -X⁵NR¹²R¹², -X⁵NR¹²C(O)R¹²,
 -X⁵NR¹²C(O)OR¹², -X⁵NR¹²C(O)NR¹²R¹², -X⁵NR¹²C(NR¹²)NR¹²R¹², -X⁵OR¹², -X⁵SR¹²,
 25 -X⁵C(O)OR¹², -X⁵C(O)R¹², -X⁵OC(O)R¹², -X⁵C(O)NR¹²R¹², -X⁵S(O)₂NR¹²R¹²,
 -X⁵NR¹²S(O)₂R¹², -X⁵P(O)(OR¹²)OR¹², -X⁵OP(O)(OR¹²)OR¹², -X⁵NR¹²C(O)R¹³, -X⁵S(O)R¹³
 and -X⁵S(O)₂R¹³ and/or 1 radical selected from -R¹⁴, -X⁵OR¹⁴, -X⁵SR¹⁴, -X⁵S(O)R¹⁴,
 -X⁵S(O)₂R¹⁴, -X⁵C(O)R¹⁴, -X⁵C(O)OR¹⁴, -X⁵OC(O)R¹⁴, -X⁵NR¹⁴R¹², -X⁵NR¹²C(O)R¹⁴,
 -X⁵NR¹²C(O)OR¹⁴, -X⁵C(O)NR¹²R¹², -X⁵S(O)₂NR¹⁴R¹², -X⁵NR¹²S(O)₂R¹⁴,
 30 -X⁵NR¹²C(O)NR¹⁴R¹² and -X⁵NR¹²C(NR¹²)NR¹⁴R¹², wherein X^5 is a bond or (C₁₋₆)alkylene;
 R¹² at each occurrence independently is hydrogen, (C₁₋₆)alkyl or halo-substituted(C₁₋₆)alkyl;
 R¹³ is (C₁₋₆)alkyl or halo-substituted(C₁₋₆)alkyl; and R¹⁴ is (C₃₋₁₀)cycloalkyl(C₀₋₆)alkyl,
 hetero(C₃₋₁₀)cycloalkyl(C₀₋₃)alkyl, (C₆₋₁₀)aryl(C₀₋₆)alkyl, hetero(C₅₋₁₀)aryl(C₀₋₆)alkyl,

(C₉₋₁₀)bicycloaryl(C₀₋₆)alkyl or hetero(C₈₋₁₀)bicycloaryl(C₀₋₆)alkyl;

R¹ is hydrogen or (C₁₋₆)alkyl and R² is selected from a group consisting of hydrogen, cyano, -X⁵NR¹²R¹², -X⁵NR¹²C(O)R¹², -X⁵NR¹²C(O)OR¹², -R¹², -X⁵NR¹²C(O)NR¹²R¹², -X⁵NR¹²C(NR¹²)NR¹²R¹², -X⁵OR¹², -X⁵SR¹², -X⁵C(O)OR¹², -X⁵C(O)R¹², -X⁵OC(O)R¹², -X⁵C(O)NR¹²R¹², -X⁵S(O)₂NR¹²R¹², -X⁵NR¹²S(O)₂R¹², -X⁵P(O)(OR¹²)OR¹², -X⁵OP(O)(OR¹²)OR¹², -X⁵NR¹²C(O)R¹³, -X⁵S(O)R¹³, -X⁵S(O)₂R¹³, -R¹⁴, -X⁵OR¹⁴, -X⁵SR¹⁴, -X⁵S(O)R¹⁴, -X⁵S(O)₂R¹⁴, -X⁵C(O)R¹⁴, -X⁵C(O)OR¹⁴, -X⁵OC(O)R¹⁴, -X⁵NR¹⁴R¹², -X⁵NR¹²C(O)R¹⁴, -X⁵NR¹²C(O)OR¹⁴, -X⁵C(O)NR¹²R¹², -X⁵S(O)₂NR¹⁴R¹², -X⁵NR¹²S(O)₂R¹⁴, -X⁵NR¹²C(O)NR¹⁴R¹² and -X⁵NR¹²C(NR¹²)NR¹⁴R¹², wherein X⁵, R¹², R¹³ and R¹⁴ are as defined above; or R¹ and R² taken together with the carbon atom to which both R¹ and R² are attached form (C₃₋₈)cycloalkylene or (C₃₋₈)heterocycloalkylene; wherein within said R² any heteroaryl, aryl, cycloalkyl, heterocycloalkyl, cycloalkylene or heterocycloalkylene is unsubstituted or substituted with 1 to 3 radicals independently selected from (C₁₋₆)alkyl, (C₁₋₆)alkylidene, cyano, halo, halo-substituted(C₁₋₄)alkyl, nitro, -X⁵NR¹²R¹², -X⁵NR¹²C(O)R¹², -X⁵NR¹²C(O)OR¹², -X⁵NR¹²C(O)NR¹²R¹², -X⁵NR¹²C(NR¹²)NR¹²R¹², -X⁵OR¹², -X⁵SR¹², -X⁵C(O)OR¹², -X⁵C(O)R¹², -X⁵OC(O)R¹², -X⁵C(O)NR¹²R¹², -X⁵S(O)₂NR¹²R¹², -X⁵NR¹²S(O)₂R¹², -X⁵P(O)(OR¹²)OR¹², -X⁵OP(O)(OR¹²)OR¹², -X⁵NR¹²C(O)R¹³, -X⁵S(O)R¹³, -X⁵S(O)₂R¹³ and -X⁵C(O)R¹³, wherein X⁵, R¹² and R¹³ are as defined above;

R³ is (C₁₋₆)alkyl or -C(R⁶)(R⁶)X⁶, wherein R⁶ is hydrogen or (C₁₋₆)alkyl and X⁶ is selected from -X⁵NR¹²R¹², -X⁵NR¹²C(O)R¹², -X⁵NR¹²C(O)OR¹², -X⁵NR¹²C(O)NR¹²R¹², -X⁵NR¹²C(NR¹²)NR¹²R¹², -X⁵OR¹², -X⁵SR¹², -X⁵C(O)OR¹², -X⁵C(O)R¹², -X⁵OC(O)R¹², -X⁵C(O)NR¹²R¹², -X⁵S(O)₂NR¹²R¹², -X⁵NR¹²S(O)₂R¹², -X⁵P(O)(OR¹²)OR¹², -X⁵OP(O)(OR¹²)OR¹², -X⁵C(O)R¹³, -X⁵NR¹²C(O)R¹³, -X⁵S(O)R¹³, -X⁵S(O)₂R¹³, -R¹⁴, -X⁵OR¹⁴, -X⁵SR¹⁴, -X⁵S(O)R¹⁴, -X⁵S(O)₂R¹⁴, -X⁵C(O)R¹⁴, -X⁵C(O)OR¹⁴, -X⁵OC(O)R¹⁴, -X⁵NR¹⁴R¹², -X⁵NR¹²C(O)R¹⁴, -X⁵NR¹²C(O)OR¹⁴, -X⁵C(O)NR¹⁴R¹², -X⁵S(O)₂NR¹⁴R¹², -X⁵NR¹²S(O)₂R¹⁴, -X⁵NR¹²C(O)NR¹⁴R¹² and -X⁵NR¹²C(NR¹²)NR¹⁴R¹² wherein X⁵, R¹², R¹³ and R¹⁴ are as defined above;

R⁴ is selected from -X⁸NR¹²R¹², -X⁸NR¹²C(O)R¹², -X⁸NR¹²C(O)OR¹², -X⁸NR¹²C(O)NR¹²R¹², -X⁸NR¹²C(NR¹²)NR¹²R¹², -X⁸OR¹², -X⁸SR¹², -X⁵C(O)OR¹², -X⁵C(O)R¹², -X⁸OC(O)R¹², -X⁵C(O)NR¹²R¹², -X⁸S(O)₂NR¹²R¹², -X⁸NR¹²S(O)₂R¹², -X⁸P(O)(OR¹²)OR¹², -X⁸OP(O)(OR¹²)OR¹², -X⁵C(O)R¹³, -X⁸NR¹²C(O)R¹³, -X⁸S(O)R¹³, -X⁸S(O)₂R¹³, -R¹⁴, -X⁸OR¹⁴, -X⁸SR¹⁴, -X⁸S(O)R¹⁴, -X⁸S(O)₂R¹⁴, -X⁵C(O)R¹⁴, -X⁵C(O)OR¹⁴, -X⁸OC(O)R¹⁴, -X⁸NR¹⁴R¹², -X⁸NR¹²C(O)R¹⁴, -X⁸NR¹²C(O)OR¹⁴, -X⁵C(O)NR¹⁴R¹²,

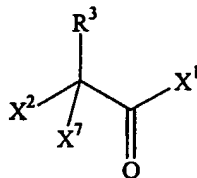
-15-

- $-X^8S(O)_2NR^{14}R^{12}$, $-X^8NR^{12}S(O)_2R^{14}$, $-X^8NR^{12}C(O)NR^{14}R^{12}$ and $-X^8NR^{12}C(NR^{12})NR^{14}R^{12}$
 wherein X^8 is (C_{1-6}) alkylene and X^5 , R^{12} , R^{13} and R^{14} are as defined above, with the proviso
 that when X^3 is cyano and X^2 is $-OR^4$, where R^4 is defined as $-R^{14}$, then R^{14} is
 (C_{3-10}) cycloalkyl (C_{1-6}) alkyl, hetero (C_{3-10}) cycloalkyl (C_{1-3}) alkyl, (C_{6-10}) aryl (C_{1-6}) alkyl,
 5 hetero (C_{5-10}) aryl (C_{1-6}) alkyl, (C_{9-10}) bicycloaryl (C_{1-6}) alkyl or
 hetero (C_{8-10}) bicycloaryl (C_{1-6}) alkyl;
 R^{15} is (C_{6-10}) aryl, hetero (C_{5-10}) aryl, (C_{9-10}) bicycloaryl or hetero (C_{8-10}) bicycloaryl;
 R^{17} is (C_{1-6}) alkyl, (C_{3-10}) cycloalkyl (C_{0-6}) alkyl, hetero (C_{3-10}) cycloalkyl (C_{0-3}) alkyl,
 (C_{6-10}) aryl (C_{0-6}) alkyl, hetero (C_{5-10}) aryl (C_{0-6}) alkyl, (C_{9-10}) bicycloaryl (C_{0-6}) alkyl or
 10 hetero (C_{8-10}) bicycloaryl (C_{0-6}) alkyl, with the proviso that when X^3 is cyano, then R^{17} is
 (C_{1-6}) alkyl, (C_{3-10}) cycloalkyl (C_{1-6}) alkyl, hetero (C_{3-10}) cycloalkyl (C_{1-6}) alkyl,
 (C_{6-10}) aryl (C_{1-6}) alkyl, hetero (C_{5-10}) aryl (C_{1-6}) alkyl, (C_{9-10}) bicycloaryl (C_{1-6}) alkyl or
 hetero (C_{8-10}) bicycloaryl (C_{1-6}) alkyl;
 R^{18} is hydrogen, (C_{1-6}) alkyl, (C_{3-10}) cycloalkyl (C_{0-6}) alkyl,
 15 hetero (C_{3-10}) cycloalkyl (C_{0-6}) alkyl, (C_{6-10}) aryl (C_{0-6}) alkyl, hetero (C_{5-10}) aryl (C_{0-6}) alkyl,
 (C_{9-10}) bicycloaryl (C_{0-6}) alkyl or hetero (C_{8-10}) bicycloaryl (C_{0-6}) alkyl, with the proviso that when
 X^3 is cyano, then R^{18} is (C_{1-6}) alkyl, (C_{3-10}) cycloalkyl (C_{1-6}) alkyl,
 hetero (C_{3-10}) cycloalkyl (C_{1-6}) alkyl, (C_{6-10}) aryl (C_{1-6}) alkyl, hetero (C_{5-10}) aryl (C_{1-6}) alkyl,
 (C_{9-10}) bicycloaryl (C_{1-6}) alkyl or hetero (C_{8-10}) bicycloaryl (C_{1-6}) alkyl; and
 20 R^{19} and R^{20} together with the atoms to which R^{19} and R^{20} are attached form
 (C_{4-8}) heterocycloalkylene, wherein no more than one of the ring member atoms comprising
 the ring is a heteroatom selected from $-NR^{21}-$ or $-O-$, wherein the ring is unsubstituted or
 substituted with R^2 , wherein R^2 is as defined above, and R^{21} is hydrogen, $-C(O)OR^{12}$,
 $-C(O)R^{12}$, $-C(O)NR^{12}R^{12}$, $-S(O)_2NR^{12}R^{12}$, $-S(O)R^{13}$ and $-S(O)_2R^{13}$, $-S(O)R^{14}$, $-S(O)_2R^{14}$,
 25 $-C(O)R^{14}$, $-C(O)OR^{14}$, $-C(O)NR^{12}R^{12}$ and $-S(O)_2NR^{14}R^{12}$, wherein R^{12} , R^{13} and R^{14} are as
 defined above;
 wherein within R^3 , R^4 , R^{15} , R^{17} and R^{18} any alicyclic or aromatic ring system is
 unsubstituted or substituted further by 1 to 5 radicals independently selected from (C_{1-6}) alkyl,
 (C_{1-6}) alkylidene, cyano, halo, halo-substituted (C_{1-4}) alkyl, nitro, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$,
 30 $-X^5NR^{12}C(O)OR^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$,
 $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$,
 $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$,
 $-X^5C(O)R^{13}$ and $-X^5S(O)_2R^{13}$ and/or 1 radical selected from $-R^{14}$, $-X^5OR^{14}$, $-X^5SR^{14}$,

-16-

$-X^5S(O)R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$, $-X^5OC(O)R^{14}$, $-X^5NR^{14}R^{12}$,
 $-X^5NR^{12}C(O)R^{14}$, $-X^5NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{14}R^{12}$, $-X^5S(O)_2NR^{14}R^{12}$, $-X^5NR^{12}S(O)_2R^{14}$,
 $-X^5NR^{12}C(O)NR^{14}R^{12}$ and $-X^5NR^{12}C(NR^{12})NR^{14}R^{12}$; and within R^3 and R^4 any aliphatic
moiety is unsubstituted or substituted further by 1 to 5 radicals independently selected from
5 cyano, halo, nitro, $-NR^{12}R^{12}$, $-NR^{12}C(O)R^{12}$, $-NR^{12}C(O)OR^{12}$, $-NR^{12}C(O)NR^{12}R^{12}$,
 $-NR^{12}C(NR^{12})NR^{12}R^{12}$, $-OR^{12}$, $-SR^{12}$, $-C(O)OR^{12}$, $-C(O)R^{12}$, $-OC(O)R^{12}$, $-C(O)NR^{12}R^{12}$,
 $-S(O)_2NR^{12}R^{12}$, $-NR^{12}S(O)_2R^{12}$, $-P(O)(OR^{12})OR^{12}$, $-OP(O)(OR^{12})OR^{12}$, $-NR^{12}C(O)R^{13}$,
 $-S(O)R^{13}$ and $-S(O)_2R^{13}$; wherein X^5 , R^{12} , R^{13} and R^{14} are as described above, with the proviso
that when X^3 is cyano and X^2 is $-OR^4$, where R^4 is defined as $-R^{14}$, or $-NHR^{18}$, then any
10 aromatic ring system present within R^{14} or R^{18} is not substituted further by halo,
 (C_{3-10}) cycloalkyl, hetero (C_{3-10}) cycloalkyl, (C_{6-10}) aryl, hetero (C_{5-10}) aryl, (C_{9-10}) bicycloaryl or
hetero (C_{8-10}) bicycloaryl; with the proviso that only one bicyclic ring structure is present
within R^3 , R^4 or R^{15} ; and the *N*-oxide derivatives, prodrug derivatives, protected derivatives,
individual isomers and mixtures of isomers thereof; and the pharmaceutically acceptable salts
15 and solvates of such compounds and the *N*-oxide derivatives, prodrug derivatives, protected
derivatives, individual isomers and mixtures of isomers thereof.

Preferred is a compound of Formula I:



I

20 in which:

X^1 is $-NHC(R^1)(R^2)X^3$ or $-NHCH(R^{19})C(O)R^{20}$;

X^2 is hydrogen, fluoro, $-OH$, $-OR^4$, $-NHR^{15}$ or $-NR^{17}R^{18}$ and X^7 is hydrogen or X^2 and
 X^7 both represent fluoro;

X^3 is $-C(R^7)(R^8)R^{16}$, $-C(R^6)(OR^6)_2$, $-CH_2C(O)R^{16}$, $-CH=CHS(O)_2R^5$,
25 $-C(O)CF_2C(O)NR^5R^5$, $-C(O)C(O)NR^5R^6$, $-C(O)C(O)OR^5$, $-C(O)CH_2OR^5$,
 $-C(O)CH_2N(R^6)SO_2R^5$ or $-C(O)C(O)R^5$; wherein R^5 is hydrogen, (C_{1-4}) alkyl,
 (C_{3-10}) cycloalkyl (C_{0-6}) alkyl, hetero (C_{3-10}) cycloalkyl (C_{0-3}) alkyl, (C_{6-10}) aryl (C_{0-6}) alkyl,
hetero (C_{5-10}) aryl (C_{0-6}) alkyl, (C_{9-10}) bicycloaryl (C_{0-6}) alkyl or
hetero (C_{8-10}) bicycloaryl (C_{0-6}) alkyl; R^6 is hydrogen, hydroxy or (C_{1-6}) alkyl; or where X^3

contains an $-NR^5R^6$ group, R^5 and R^6 together with the nitrogen atom to which they are both attached, form hetero(C₃₋₁₀)cycloalkyl, hetero(C₅₋₁₀)aryl or hetero(C₈₋₁₀)bicycloaryl; R^7 is hydrogen or (C₁₋₄)alkyl and R^8 is hydroxy or R^7 and R^8 together form oxo; R^{16} is hydrogen, $-X^4$, $-CF_3$, $-CF_2CF_2R^9$ or $-N(R^6)OR^6$; R^9 is hydrogen, halo, (C₁₋₄)alkyl, (C₅₋₁₀)aryl(C₀₋₆)alkyl or (C₅₋₁₀)heteroaryl(C₀₋₆)alkyl;

X^4 comprises a heteromonocyclic ring containing 4 to 7 ring member atoms or a fused heterobicyclic ring system containing 8 to 14 ring member atoms and any carbocyclic ketone, iminoketone or thioketone derivative thereof, with the proviso that when $-X^4$ is other than a heteromonocyclic ring containing 5 ring member atoms, wherein no more than two of the ring member atoms comprising the ring are heteroatoms, then X^2 is fluoro, $-OH$, $-OR^4$, $-NHR^{15}$ or $-NR^{17}R^{18}$ and X^7 is hydrogen or X^2 and X^7 both represent fluoro;

wherein within R^5 , X^3 or X^4 any alicyclic or aromatic ring system is unsubstituted or substituted further by 1 to 5 radicals independently selected from (C₁₋₆)alkyl, (C₁₋₆)alkylidene, cyano, halo, halo-substituted(C₁₋₄)alkyl, nitro, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$, $-X^5NR^{12}C(O)OR^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$, $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$, $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$ and $-X^5S(O)_2R^{13}$ and/or 1 radical selected from $-R^{14}$, $-X^5OR^{14}$, $-X^5SR^{14}$, $-X^5S(O)R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$, $-X^5OC(O)R^{14}$, $-X^5NR^{14}R^{12}$, $-X^5NR^{12}C(O)R^{14}$, $-X^5NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{14}R^{12}$, $-X^5NR^{12}S(O)_2R^{14}$, $-X^5NR^{12}C(O)NR^{14}R^{12}$ and $-X^5NR^{12}C(NR^{12})NR^{14}R^{12}$, wherein X^5 is a bond or (C₁₋₆)alkylene; R^{12} at each occurrence independently is hydrogen, (C₁₋₆)alkyl or halo-substituted(C₁₋₆)alkyl; R^{13} is (C₁₋₆)alkyl or halo-substituted(C₁₋₆)alkyl; and R^{14} is (C₃₋₁₀)cycloalkyl(C₀₋₆)alkyl, hetero(C₃₋₁₀)cycloalkyl(C₀₋₃)alkyl, (C₆₋₁₀)aryl(C₀₋₆)alkyl, hetero(C₅₋₁₀)aryl(C₀₋₆)alkyl, (C₉₋₁₀)bicycloaryl(C₀₋₆)alkyl or hetero(C₈₋₁₀)bicycloaryl(C₀₋₆)alkyl;

R^1 is hydrogen or (C₁₋₆)alkyl and R^2 is selected from a group consisting of hydrogen, cyano, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$, $-X^5NR^{12}C(O)OR^{12}$, $-X^5R^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$, $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$, $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$, $-X^5S(O)_2R^{13}$, $-R^{14}$, $-X^5OR^{14}$, $-X^5SR^{14}$, $-X^5S(O)R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$, $-X^5OC(O)R^{14}$, $-X^5NR^{14}R^{12}$, $-X^5NR^{12}C(O)R^{14}$, $-X^5NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{14}R^{12}$, $-X^5NR^{12}S(O)_2R^{14}$, $-X^5NR^{12}C(O)NR^{14}R^{12}$ and $-X^5NR^{12}C(NR^{12})NR^{14}R^{12}$, wherein X^5 , R^{12} , R^{13} and R^{14} are as

-18-

defined above; or R^1 and R^2 taken together with the carbon atom to which both R^1 and R^2 are attached form (C_{3-8}) cycloalkylene or (C_{3-8}) heterocycloalkylene; wherein within said R^2 any heteroaryl, aryl, cycloalkyl, heterocycloalkyl, cycloalkylene or heterocycloalkylene is unsubstituted or substituted with 1 to 3 radicals independently selected from (C_{1-6}) alkyl,

5 (C_{1-6}) alkylidene, cyano, halo, halo-substituted (C_{1-4}) alkyl, nitro, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$, $-X^5NR^{12}C(O)OR^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$, $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$, $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$, $-X^5S(O)_2R^{13}$ and $-X^5C(O)R^{13}$, wherein X^5 , R^{12} and R^{13} are as defined above;

10 R^3 is $-C(R^6)(R^6)X^6$, wherein R^6 is hydrogen or (C_{1-6}) alkyl and X^6 is selected from $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$, $-X^5NR^{12}C(O)OR^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$, $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$, $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5C(O)R^{13}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$, $-X^5S(O)_2R^{13}$, $-R^{14}$, $-X^5OR^{14}$, $-X^5SR^{14}$, $-X^5S(O)R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$, $-X^5OC(O)R^{14}$, $-X^5NR^{14}R^{12}$, $-X^5NR^{12}C(O)R^{14}$, $-X^5NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{14}R^{12}$, $-X^5S(O)_2NR^{14}R^{12}$, $-X^5NR^{12}S(O)_2R^{14}$, $-X^5NR^{12}C(O)NR^{14}R^{12}$ and $-X^5NR^{12}C(NR^{12})NR^{14}R^{12}$ wherein X^5 , R^{12} , R^{13} and R^{14} are as defined above;

R^4 is selected from $-X^8NR^{12}R^{12}$, $-X^8NR^{12}C(O)R^{12}$, $-X^8NR^{12}C(O)OR^{12}$, $-X^8NR^{12}C(O)NR^{12}R^{12}$, $-X^8NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^8OR^{12}$, $-X^8SR^{12}$, $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^8OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^8S(O)_2NR^{12}R^{12}$, $-X^8NR^{12}S(O)_2R^{12}$, $-X^8P(O)(OR^{12})OR^{12}$, $-X^8OP(O)(OR^{12})OR^{12}$, $-X^5C(O)R^{13}$, $-X^8NR^{12}C(O)R^{13}$, $-X^8S(O)R^{13}$, $-X^8S(O)_2R^{13}$, $-R^{14}$, $-X^8OR^{14}$, $-X^8SR^{14}$, $-X^8S(O)R^{14}$, $-X^8S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$, $-X^8OC(O)R^{14}$, $-X^8NR^{14}R^{12}$, $-X^8NR^{12}C(O)R^{14}$, $-X^8NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{14}R^{12}$, $-X^8S(O)_2NR^{14}R^{12}$, $-X^8NR^{12}S(O)_2R^{14}$, $-X^8NR^{12}C(O)NR^{14}R^{12}$ and $-X^8NR^{12}C(NR^{12})NR^{14}R^{12}$ wherein X^8 is (C_{1-6}) alkylene and X^5 , R^{12} , R^{13} and R^{14} are as defined above;

R^{15} is (C_{6-10}) aryl, hetero (C_{5-10}) aryl, (C_{9-10}) bicycloaryl or hetero (C_{8-10}) bicycloaryl;

R^{17} is hydrogen, (C_{1-6}) alkyl, (C_{3-10}) cycloalkyl (C_{0-6}) alkyl, hetero (C_{3-10}) cycloalkyl (C_{0-3}) alkyl, (C_{6-10}) aryl (C_{0-6}) alkyl, hetero (C_{5-10}) aryl (C_{0-6}) alkyl, (C_{9-10}) bicycloaryl (C_{0-6}) alkyl or hetero (C_{8-10}) bicycloaryl (C_{0-6}) alkyl;

R^{18} is (C_{1-6}) alkyl, (C_{3-10}) cycloalkyl (C_{0-6}) alkyl, hetero (C_{3-10}) cycloalkyl (C_{0-6}) alkyl, (C_{6-10}) aryl (C_{0-6}) alkyl, hetero (C_{5-10}) aryl (C_{0-6}) alkyl, (C_{9-10}) bicycloaryl (C_{0-6}) alkyl or hetero (C_{8-10}) bicycloaryl (C_{0-6}) alkyl; and

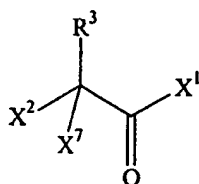
-19-

R^{19} and R^{20} together with the atoms to which R^{19} and R^{20} are attached form (C₄₋₈)heterocycloalkylene, wherein no more than one of the ring member atoms comprising the ring is a heteroatom selected from -NR²¹- or -O-, wherein the ring is unsubstituted or substituted with R^1 , wherein R^1 is as defined above, and R^{21} is hydrogen, -C(O)OR¹²,
 5 -C(O)R¹², -C(O)NR¹²R¹², -S(O)₂NR¹²R¹², -S(O)R¹³ and -S(O)₂R¹³, -S(O)R¹⁴, -S(O)₂R¹⁴,
 -C(O)R¹⁴, -C(O)OR¹⁴, -C(O)NR¹²R¹² and -S(O)₂NR¹⁴R¹², wherein R^{12} , R^{13} and R^{14} are as defined above;

wherein within R^3 , R^4 , R^{15} , R^{17} and R^{18} any alicyclic or aromatic ring system is unsubstituted or substituted further by 1 to 5 radicals independently selected from (C₁₋₆)alkyl,
 10 (C₁₋₆)alkylidene, cyano, halo, halo-substituted(C₁₋₄)alkyl, nitro, -X⁵NR¹²R¹², -X⁵NR¹²C(O)R¹²,
 -X⁵NR¹²C(O)OR¹², -X⁵NR¹²C(O)NR¹²R¹², -X⁵NR¹²C(NR¹²)NR¹²R¹², -X⁵OR¹², -X⁵SR¹²,
 -X⁵C(O)OR¹², -X⁵C(O)R¹², -X⁵OC(O)R¹², -X⁵C(O)NR¹²R¹², -X⁵S(O)₂NR¹²R¹²,
 -X⁵NR¹²S(O)₂R¹², -X⁵P(O)(OR¹²)OR¹², -X⁵OP(O)(OR¹²)OR¹², -X⁵NR¹²C(O)R¹³, -X⁵S(O)R¹³,
 -X⁵C(O)R¹³ and -X⁵S(O)₂R¹³ and/or 1 radical selected from -R¹⁴, -X⁵OR¹⁴, -X⁵SR¹⁴,
 15 -X⁵S(O)R¹⁴, -X⁵S(O)₂R¹⁴, -X⁵C(O)R¹⁴, -X⁵C(O)OR¹⁴, -X⁵OC(O)R¹⁴, -X⁵NR¹⁴R¹²,
 -X⁵NR¹²C(O)R¹⁴, -X⁵NR¹²C(O)OR¹⁴, -X⁵C(O)NR¹⁴R¹², -X⁵S(O)₂NR¹⁴R¹², -X⁵NR¹²S(O)₂R¹⁴,
 -X⁵NR¹²C(O)NR¹⁴R¹² and -X⁵NR¹²C(NR¹²)NR¹⁴R¹²; and within R^3 and R^4 any aliphatic moiety is unsubstituted or substituted further by 1 to 5 radicals independently selected from
 cyano, halo, nitro, -NR¹²R¹², -NR¹²C(O)R¹², -NR¹²C(O)OR¹², -NR¹²C(O)NR¹²R¹²,
 20 -NR¹²C(NR¹²)NR¹²R¹², -OR¹², -SR¹², -C(O)OR¹², -C(O)R¹², -OC(O)R¹², -C(O)NR¹²R¹²,
 -S(O)₂NR¹²R¹², -NR¹²S(O)₂R¹², -P(O)(OR¹²)OR¹², -OP(O)(OR¹²)OR¹², -NR¹²C(O)R¹³,
 -S(O)R¹³ and -S(O)₂R¹³; wherein X⁵, R^{12} , R^{13} and R^{14} are as described above; with the proviso that only one bicyclic ring structure is present within R^3 , R^4 or R^{15} ; and the *N*-oxide derivatives, prodrug derivatives, protected derivatives, individual isomers and mixtures of
 25 isomers thereof; and the pharmaceutically acceptable salts and solvates of such compounds and the *N*-oxide derivatives, prodrug derivatives, protected derivatives, individual isomers and mixtures of isomers thereof.

Preferred is a compound of Formula I:

-20-



I

in which:

X^1 is $-NHC(R^1)(R^2)X^3$ or $-NHCH(R^{19})C(O)R^{20}$;

X^2 is hydrogen, fluoro, $-OH$, $-OR^4$ or $-NR^{17}R^{18}$ and X^7 is hydrogen or X^2 and X^7 both
 5 represent fluoro;

X^3 is cyano;

wherein within X^3 any alicyclic or aromatic ring system is unsubstituted or substituted
 further by 1 to 5 radicals independently selected from (C_{1-6}) alkyl, (C_{1-6}) alkylidene, cyano,
 halo, halo-substituted (C_{1-4}) alkyl, nitro, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$, $-X^5NR^{12}C(O)OR^{12}$,
 10 $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$, $-X^5C(O)OR^{12}$,
 $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$, $-X^5NR^{12}S(O)_2R^{12}$,
 $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$ and $-X^5S(O)_2R^{13}$
 and/or 1 radical selected from $-R^{14}$, $-X^5OR^{14}$, $-X^5SR^{14}$, $-X^5S(O)R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$,
 $-X^5C(O)OR^{14}$, $-X^5OC(O)R^{14}$, $-X^5NR^{14}R^{12}$, $-X^5NR^{12}C(O)R^{14}$, $-X^5NR^{12}C(O)OR^{14}$,
 15 $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{14}R^{12}$, $-X^5NR^{12}S(O)_2R^{14}$, $-X^5NR^{12}C(O)NR^{14}R^{12}$ and
 $-X^5NR^{12}C(NR^{12})NR^{14}R^{12}$, wherein X^5 is a bond or (C_{1-6}) alkylene; R^{12} at each occurrence
 independently is hydrogen, (C_{1-6}) alkyl or halo-substituted (C_{1-6}) alkyl; R^{13} is (C_{1-6}) alkyl or
 halo-substituted (C_{1-6}) alkyl; and R^{14} is (C_{3-10}) cycloalkyl (C_{0-6}) alkyl,
 hetero (C_{3-10}) cycloalkyl (C_{0-3}) alkyl, (C_{6-10}) aryl (C_{0-6}) alkyl, hetero (C_{5-10}) aryl (C_{0-6}) alkyl,
 20 (C_{9-10}) bicycloaryl (C_{0-6}) alkyl or hetero (C_{8-10}) bicycloaryl (C_{0-6}) alkyl;

R^1 is hydrogen or (C_{1-6}) alkyl and R^2 is selected from a group consisting of hydrogen,
 cyano, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$, $-X^5NR^{12}C(O)OR^{12}$, $-X^5R^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$,
 $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$, $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$,
 $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$, $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$,
 25 $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$, $-X^5S(O)_2R^{13}$, $-R^{14}$, $-X^5OR^{14}$, $-X^5SR^{14}$,
 $-X^5S(O)R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$, $-X^5OC(O)R^{14}$, $-X^5NR^{14}R^{12}$,
 $-X^5NR^{12}C(O)R^{14}$, $-X^5NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{14}R^{12}$, $-X^5NR^{12}S(O)_2R^{14}$,
 $-X^5NR^{12}C(O)NR^{14}R^{12}$ and $-X^5NR^{12}C(NR^{12})NR^{14}R^{12}$, wherein X^5 , R^{12} , R^{13} and R^{14} are as
 defined above; or R^1 and R^2 taken together with the carbon atom to which both R^1 and R^2 are

attached form (C₃₋₈)cycloalkylene or (C₃₋₈)heterocycloalkylene; wherein within said R² any heteroaryl, aryl, cycloalkyl, heterocycloalkyl, cycloalkylene or heterocycloalkylene is unsubstituted or substituted with 1 to 3 radicals independently selected from (C₁₋₆)alkyl, (C₁₋₆)alkylidene, cyano, halo, halo-substituted(C₁₋₄)alkyl, nitro, -X⁵NR¹²R¹², -X⁵NR¹²C(O)R¹²,
 5 -X⁵NR¹²C(O)OR¹², -X⁵NR¹²C(O)NR¹²R¹², -X⁵NR¹²C(NR¹²)NR¹²R¹², -X⁵OR¹², -X⁵SR¹²,
 -X⁵C(O)OR¹², -X⁵C(O)R¹², -X⁵OC(O)R¹², -X⁵C(O)NR¹²R¹², -X⁵S(O)₂NR¹²R¹²,
 -X⁵NR¹²S(O)₂R¹², -X⁵P(O)(OR¹²)OR¹², -X⁵OP(O)(OR¹²)OR¹², -X⁵NR¹²C(O)R¹³, -X⁵S(O)R¹³,
 -X⁵S(O)₂R¹³ and -X⁵C(O)R¹³, wherein X⁵, R¹² and R¹³ are as defined above;

R³ is -C(R⁶)(R⁶)X⁶, wherein R⁶ is hydrogen or (C₁₋₆)alkyl and X⁶ is selected from
 10 -X⁵NR¹²R¹², -X⁵NR¹²C(O)R¹², -X⁵NR¹²C(O)OR¹², -X⁵NR¹²C(O)NR¹²R¹²,
 -X⁵NR¹²C(NR¹²)NR¹²R¹², -X⁵OR¹², -X⁵SR¹², -X⁵C(O)OR¹², -X⁵C(O)R¹², -X⁵OC(O)R¹²,
 -X⁵C(O)NR¹²R¹², -X⁵S(O)₂NR¹²R¹², -X⁵NR¹²S(O)₂R¹², -X⁵P(O)(OR¹²)OR¹²,
 -X⁵OP(O)(OR¹²)OR¹², -X⁵C(O)R¹³, -X⁵NR¹²C(O)R¹³, -X⁵S(O)R¹³, -X⁵S(O)₂R¹³, -R¹⁴,
 -X⁵OR¹⁴, -X⁵SR¹⁴, -X⁵S(O)R¹⁴, -X⁵S(O)₂R¹⁴, -X⁵C(O)R¹⁴, -X⁵C(O)OR¹⁴, -X⁵OC(O)R¹⁴,
 15 -X⁵NR¹⁴R¹², -X⁵NR¹²C(O)R¹⁴, -X⁵NR¹²C(O)OR¹⁴, -X⁵C(O)NR¹⁴R¹², -X⁵S(O)₂NR¹⁴R¹²,
 -X⁵NR¹²S(O)₂R¹⁴, -X⁵NR¹²C(O)NR¹⁴R¹² and -X⁵NR¹²C(NR¹²)NR¹⁴R¹² wherein X⁵, R¹², R¹³
 and R¹⁴ are as defined above;

R⁴ is selected from -X⁸NR¹²R¹², -X⁸NR¹²C(O)R¹², -X⁸NR¹²C(O)OR¹²,
 -X⁸NR¹²C(O)NR¹²R¹², -X⁸NR¹²C(NR¹²)NR¹²R¹², -X⁸OR¹², -X⁸SR¹², -X⁵C(O)OR¹²,
 20 -X⁵C(O)R¹², -X⁸OC(O)R¹², -X⁵C(O)NR¹²R¹², -X⁸S(O)₂NR¹²R¹², -X⁸NR¹²S(O)₂R¹²,
 -X⁸P(O)(OR¹²)OR¹², -X⁸OP(O)(OR¹²)OR¹², -X⁵C(O)R¹³, -X⁸NR¹²C(O)R¹³, -X⁸S(O)R¹³,
 -X⁸S(O)₂R¹³, -R¹⁴, -X⁸OR¹⁴, -X⁸SR¹⁴, -X⁸S(O)R¹⁴, -X⁸S(O)₂R¹⁴, -X⁵C(O)R¹⁴, -X⁵C(O)OR¹⁴,
 -X⁸OC(O)R¹⁴, -X⁸NR¹⁴R¹², -X⁸NR¹²C(O)R¹⁴, -X⁸NR¹²C(O)OR¹⁴, -X⁵C(O)NR¹⁴R¹²,
 -X⁸S(O)₂NR¹⁴R¹², -X⁸NR¹²S(O)₂R¹⁴, -X⁸NR¹²C(O)NR¹⁴R¹² and -X⁸NR¹²C(NR¹²)NR¹⁴R¹²
 25 wherein X⁸ is (C₁₋₆)alkylene and X⁵, R¹², R¹³ and R¹⁴ are as defined above, with the proviso
 that when X³ is cyano and X² is -OR⁴, where R⁴ is defined as -R¹⁴, then R¹⁴ is
 (C₃₋₁₀)cycloalkyl(C₁₋₆)alkyl, hetero(C₃₋₁₀)cycloalkyl(C₁₋₃)alkyl, (C₆₋₁₀)aryl(C₁₋₆)alkyl,
 hetero(C₅₋₁₀)aryl(C₁₋₆)alkyl, (C₉₋₁₀)bicycloaryl(C₁₋₆)alkyl or
 hetero(C₈₋₁₀)bicycloaryl(C₁₋₆)alkyl;

30 R¹⁵ is (C₆₋₁₀)aryl, hetero(C₅₋₁₀)aryl, (C₉₋₁₀)bicycloaryl or hetero(C₈₋₁₀)bicycloaryl;

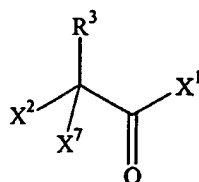
R¹⁷ is (C₁₋₆)alkyl, (C₃₋₁₀)cycloalkyl(C₁₋₆)alkyl, hetero(C₃₋₁₀)cycloalkyl(C₁₋₆)alkyl,
 (C₆₋₁₀)aryl(C₁₋₆)alkyl, hetero(C₅₋₁₀)aryl(C₁₋₆)alkyl, (C₉₋₁₀)bicycloaryl(C₁₋₆)alkyl or
 hetero(C₈₋₁₀)bicycloaryl(C₁₋₆)alkyl;

R^{18} is (C_{1-6}) alkyl, (C_{3-10}) cycloalkyl (C_{1-6}) alkyl, hetero (C_{3-10}) cycloalkyl (C_{1-6}) alkyl, (C_{6-10}) aryl (C_{1-6}) alkyl, hetero (C_{5-10}) aryl (C_{1-6}) alkyl, (C_{9-10}) bicycloaryl (C_{1-6}) alkyl or hetero (C_{8-10}) bicycloaryl (C_{1-6}) alkyl; and

- R^{19} and R^{20} together with the atoms to which R^{19} and R^{20} are attached form
- 5 (C_{4-8}) heterocycloalkylene, wherein no more than one of the ring member atoms comprising the ring is a heteroatom selected from $-NR^{21}-$ or $-O-$, wherein the ring is unsubstituted or substituted with R^1 , wherein R^1 is as defined above, and R^{21} is hydrogen, $-C(O)OR^{12}$, $-C(O)R^{12}$, $-C(O)NR^{12}R^{12}$, $-S(O)_2NR^{12}R^{12}$, $-S(O)R^{13}$ and $-S(O)_2R^{13}$, $-S(O)R^{14}$, $-S(O)_2R^{14}$, $-C(O)R^{14}$, $-C(O)OR^{14}$, $-C(O)NR^{12}R^{12}$ and $-S(O)_2NR^{14}R^{12}$, wherein R^{12} , R^{13} and R^{14} are as
- 10 defined above;
- wherein within R^3 , R^4 , R^{15} , R^{17} and R^{18} any alicyclic or aromatic ring system is unsubstituted or substituted further by 1 to 5 radicals independently selected from (C_{1-6}) alkyl, (C_{1-6}) alkylidene, cyano, halo, halo-substituted (C_{1-4}) alkyl, nitro, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$, $-X^5NR^{12}C(O)OR^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$,
- 15 $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$, $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$, $-X^5C(O)R^{13}$ and $-X^5S(O)_2R^{13}$ and/or 1 radical selected from $-R^{14}$, $-X^5OR^{14}$, $-X^5SR^{14}$, $-X^5S(O)R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$, $-X^5OC(O)R^{14}$, $-X^5NR^{14}R^{12}$, $-X^5NR^{12}C(O)R^{14}$, $-X^5NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{14}R^{12}$, $-X^5S(O)_2NR^{14}R^{12}$, $-X^5NR^{12}S(O)_2R^{14}$,
- 20 $-X^5NR^{12}C(O)NR^{14}R^{12}$ and $-X^5NR^{12}C(NR^{12})NR^{14}R^{12}$; and within R^3 and R^4 any aliphatic moiety is unsubstituted or substituted further by 1 to 5 radicals independently selected from cyano, halo, nitro, $-NR^{12}R^{12}$, $-NR^{12}C(O)R^{12}$, $-NR^{12}C(O)OR^{12}$, $-NR^{12}C(O)NR^{12}R^{12}$, $-NR^{12}C(NR^{12})NR^{12}R^{12}$, $-OR^{12}$, $-SR^{12}$, $-C(O)OR^{12}$, $-C(O)R^{12}$, $-OC(O)R^{12}$, $-C(O)NR^{12}R^{12}$, $-S(O)_2NR^{12}R^{12}$, $-NR^{12}S(O)_2R^{12}$, $-P(O)(OR^{12})OR^{12}$, $-OP(O)(OR^{12})OR^{12}$, $-NR^{12}C(O)R^{13}$,
- 25 $-S(O)R^{13}$ and $-S(O)_2R^{13}$; wherein X^5 , R^{12} , R^{13} and R^{14} are as described above, with the proviso that when X^2 is $-OR^4$, where R^4 is defined as $-R^{14}$, or $-NHR^{18}$, then any aromatic ring system present within R^{14} or R^{18} is not substituted further by halo, (C_{3-10}) cycloalkyl, hetero (C_{3-10}) cycloalkyl, (C_{6-10}) aryl, hetero (C_{5-10}) aryl, (C_{9-10}) bicycloaryl or hetero (C_{8-10}) bicycloaryl; with the proviso that only one bicyclic ring structure is present
- 30 within R^3 , R^4 or R^{15} ; and the *N*-oxide derivatives, prodrug derivatives, protected derivatives, individual isomers and mixtures of isomers thereof; and the pharmaceutically acceptable salts and solvates of such compounds and the *N*-oxide derivatives, prodrug derivatives, protected derivatives, individual isomers and mixtures of isomers thereof.

-23-

Preferred is a compound of Formula I:



I

in which:

- 5 X^1 is $-NHC(R^1)(R^2)X^3$ or $-NHCH(R^{19})C(O)R^{20}$;
 X^2 is $-OH$, $-OC(O)NR^{12}R^{12}$ or $-OC(O)R^{14}$, wherein R^{12} and R^{14} are as defined below;
 X^3 is cyano, $-C(R^7)(R^8)R^{16}$, $-C(R^6)(OR^6)_2$, $-CH_2C(O)R^{16}$, $-CH=CHS(O)_2R^5$,
 $-C(O)CF_2C(O)NR^5R^5$, $-C(O)C(O)NR^5R^6$, $-C(O)C(O)OR^5$, $-C(O)CH_2OR^5$,
 $-C(O)CH_2N(R^6)SO_2R^5$ or $-C(O)C(O)R^5$; wherein R^5 is hydrogen, (C_{1-4}) alkyl,
10 (C_{3-10}) cycloalkyl (C_{0-6}) alkyl, hetero (C_{3-10}) cycloalkyl (C_{0-3}) alkyl, (C_{6-10}) aryl (C_{0-6}) alkyl,
hetero (C_{5-10}) aryl (C_{0-6}) alkyl, (C_{9-10}) bicycloaryl (C_{0-6}) alkyl or
hetero (C_{8-10}) bicycloaryl (C_{0-6}) alkyl; R^6 is hydrogen, hydroxy or (C_{1-6}) alkyl; or where X^3
contains an $-NR^5R^6$ group, R^5 and R^6 together with the nitrogen atom to which they are both
attached, form hetero (C_{3-10}) cycloalkyl, hetero (C_{5-10}) aryl or hetero (C_{8-10}) bicycloaryl; R^7 is
15 hydrogen or (C_{1-4}) alkyl and R^8 is hydroxy or R^7 and R^8 together form oxo; R^{16} is hydrogen, -
 X^4 , $-CF_3$, $-CF_2CF_2R^9$ or $-N(R^6)OR^6$; R^9 is hydrogen, halo, (C_{1-4}) alkyl, (C_{5-10}) aryl (C_{0-6}) alkyl or
 (C_{5-10}) heteroaryl (C_{0-6}) alkyl;

20 X^4 comprises a heteromonocyclic ring containing 4 to 7 ring member atoms or a fused
heterobicyclic ring system containing 8 to 14 ring member atoms and any carbocyclic ketone,
iminoketone or thioketone derivative thereof;

wherein within R^5 , X^3 or X^4 any alicyclic or aromatic ring system is unsubstituted or
substituted further by 1 to 5 radicals independently selected from (C_{1-6}) alkyl, (C_{1-6}) alkylidene,
cyano, halo, halo-substituted (C_{1-4}) alkyl, nitro, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$,
 $-X^5NR^{12}C(O)OR^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$,
25 $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$,
 $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$
and $-X^5S(O)_2R^{13}$ and/or 1 radical selected from $-R^{14}$, $-X^5OR^{14}$, $-X^5SR^{14}$, $-X^5S(O)R^{14}$,
 $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$, $-X^5OC(O)R^{14}$, $-X^5NR^{14}R^{12}$, $-X^5NR^{12}C(O)R^{14}$,
 $-X^5NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{14}R^{12}$, $-X^5NR^{12}S(O)_2R^{14}$,

$-X^5NR^{12}C(O)NR^{14}R^{12}$ and $-X^5NR^{12}C(NR^{12})NR^{14}R^{12}$, wherein X^5 is a bond or (C_{1-6}) alkylene; R^{12} at each occurrence independently is hydrogen, (C_{1-6}) alkyl or halo-substituted (C_{1-6}) alkyl; R^{13} is (C_{1-6}) alkyl or halo-substituted (C_{1-6}) alkyl; and R^{14} is (C_{3-10}) cycloalkyl (C_{0-6}) alkyl, hetero (C_{3-10}) cycloalkyl (C_{0-3}) alkyl, (C_{6-10}) aryl (C_{0-6}) alkyl, hetero (C_{5-10}) aryl (C_{0-6}) alkyl,
 5 (C_{9-10}) bicycloaryl (C_{0-6}) alkyl or hetero (C_{8-10}) bicycloaryl (C_{0-6}) alkyl;

R^1 is hydrogen or (C_{1-6}) alkyl and R^2 is selected from a group consisting of hydrogen, cyano, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$, $-X^5NR^{12}C(O)OR^{12}$, $-X^5R^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$, $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$, $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$,
 10 $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$, $-X^5S(O)_2R^{13}$, $-R^{14}$, $-X^5OR^{14}$, $-X^5SR^{14}$, $-X^5S(O)R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$, $-X^5OC(O)R^{14}$, $-X^5NR^{14}R^{12}$, $-X^5NR^{12}C(O)R^{14}$, $-X^5NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{14}R^{12}$, $-X^5NR^{12}S(O)_2R^{14}$, $-X^5NR^{12}C(O)NR^{14}R^{12}$ and $-X^5NR^{12}C(NR^{12})NR^{14}R^{12}$, wherein X^5 , R^{12} , R^{13} and R^{14} are as defined above; or R^1 and R^2 taken together with the carbon atom to which both R^1 and R^2 are
 15 attached form (C_{3-8}) cycloalkylene or (C_{3-8}) heterocycloalkylene; wherein within said R^2 any heteroaryl, aryl, cycloalkyl, heterocycloalkyl, cycloalkylene or heterocycloalkylene is unsubstituted or substituted with 1 to 3 radicals independently selected from (C_{1-6}) alkyl, (C_{1-6}) alkylidene, cyano, halo, halo-substituted (C_{1-4}) alkyl, nitro, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$, $-X^5NR^{12}C(O)OR^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$,
 20 $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$, $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$, $-X^5S(O)_2R^{13}$ and $-X^5C(O)R^{13}$, wherein X^5 , R^{12} and R^{13} are as defined above;

R^3 is $-C(R^6)(R^6)X^6$, wherein R^6 is hydrogen or (C_{1-6}) alkyl and X^6 is selected from $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$, $-X^5NR^{12}C(O)OR^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$,
 25 $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$, $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$, $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$, $-X^5S(O)_2R^{13}$, $-R^{14}$, $-X^5OR^{14}$, $-X^5SR^{14}$, $-X^5S(O)R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$, $-X^5OC(O)R^{14}$, $-X^5NR^{14}R^{12}$, $-X^5NR^{12}C(O)R^{14}$, $-X^5NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{14}R^{12}$, $-X^5S(O)_2NR^{14}R^{12}$,
 30 $-X^5NR^{12}S(O)_2R^{14}$, $-X^5NR^{12}C(O)NR^{14}R^{12}$ and $-X^5NR^{12}C(NR^{12})NR^{14}R^{12}$ wherein X^5 , R^{12} , R^{13} and R^{14} are as defined above; and

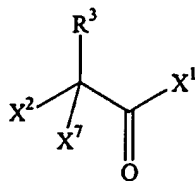
R^{19} and R^{20} together with the atoms to which R^{19} and R^{20} are attached form (C_{4-8}) heterocycloalkylene, wherein no more than one of the ring member atoms comprising

-25-

the ring is a heteroatom selected from $-NR^{21}-$ or $-O-$, wherein and the ring is unsubstituted or substituted with R^1 , wherein R^1 is as defined above, and R^{21} is hydrogen, $-C(O)OR^{12}$, $-C(O)R^{12}$, $-C(O)NR^{12}R^{12}$, $-S(O)_2NR^{12}R^{12}$, $-S(O)R^{13}$ and $-S(O)_2R^{13}$, $-S(O)R^{14}$, $-S(O)_2R^{14}$, $-C(O)R^{14}$, $-C(O)OR^{14}$, $-C(O)NR^{12}R^{12}$ and $-S(O)_2NR^{14}R^{12}$, wherein R^{12} , R^{13} and R^{14} are as defined above;

wherein within R^3 , R^4 , R^{15} , R^{17} and R^{18} any alicyclic or aromatic ring system is unsubstituted or substituted further by 1 to 5 radicals independently selected from (C_{1-6}) alkyl, (C_{1-6}) alkylidene, cyano, halo, halo-substituted (C_{1-4}) alkyl, nitro, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$, $-X^5NR^{12}C(O)OR^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$, $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$, $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$, $-X^5C(O)R^{13}$ and $-X^5S(O)_2R^{13}$ and/or 1 radical selected from $-R^{14}$, $-X^5OR^{14}$, $-X^5SR^{14}$, $-X^5S(O)R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$, $-X^5OC(O)R^{14}$, $-X^5NR^{14}R^{12}$, $-X^5NR^{12}C(O)R^{14}$, $-X^5NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{14}R^{12}$, $-X^5S(O)_2NR^{14}R^{12}$, $-X^5NR^{12}S(O)_2R^{14}$, $-X^5NR^{12}C(O)NR^{14}R^{12}$ and $-X^5NR^{12}C(NR^{12})NR^{14}R^{12}$; and within R^3 and R^4 any aliphatic moiety is unsubstituted or substituted further by 1 to 5 radicals independently selected from cyano, halo, nitro, $-NR^{12}R^{12}$, $-NR^{12}C(O)R^{12}$, $-NR^{12}C(O)OR^{12}$, $-NR^{12}C(O)NR^{12}R^{12}$, $-NR^{12}C(NR^{12})NR^{12}R^{12}$, $-OR^{12}$, $-SR^{12}$, $-C(O)OR^{12}$, $-C(O)R^{12}$, $-OC(O)R^{12}$, $-C(O)NR^{12}R^{12}$, $-S(O)_2NR^{12}R^{12}$, $-NR^{12}S(O)_2R^{12}$, $-P(O)(OR^{12})OR^{12}$, $-OP(O)(OR^{12})OR^{12}$, $-NR^{12}C(O)R^{13}$, $-S(O)R^{13}$ and $-S(O)_2R^{13}$; wherein X^5 , R^{12} , R^{13} and R^{14} are as described above; with the proviso that only one bicyclic ring structure is present within R^3 , R^4 or R^{15} ; and the *N*-oxide derivatives, prodrug derivatives, protected derivatives, individual isomers and mixtures of isomers thereof; and the pharmaceutically acceptable salts and solvates of such compounds and the *N*-oxide derivatives, prodrug derivatives, protected derivatives, individual isomers and mixtures of isomers thereof.

Preferred is a compound of Formula I:



I

in which:

X^1 is $-NHC(R^1)(R^2)C(O)C(O)NR^5R^6$, wherein R^5 is hydrogen, (C_{1-4}) alkyl, (C_{3-10}) cycloalkyl (C_{0-6}) alkyl, hetero (C_{3-10}) cycloalkyl (C_{0-3}) alkyl, (C_{6-10}) aryl (C_{0-6}) alkyl, hetero (C_{5-10}) aryl (C_{0-6}) alkyl, (C_{9-10}) bicycloaryl (C_{0-6}) alkyl or hetero (C_{8-10}) bicycloaryl (C_{0-6}) alkyl and R^6 is hydrogen, hydroxy or (C_{1-6}) alkyl or R^5 and R^6 together with the nitrogen atom to which they are both attached form hetero (C_{3-10}) cycloalkyl, hetero (C_{5-10}) aryl or hetero (C_{8-10}) bicycloaryl;

X^2 is hydrogen;

wherein within X^1 any alicyclic or aromatic ring system is unsubstituted or substituted further by 1 to 5 radicals independently selected from (C_{1-6}) alkyl, (C_{1-6}) alkylidene, cyano, halo, halo-substituted (C_{1-4}) alkyl, nitro, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$, $-X^5NR^{12}C(O)OR^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$, $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$, $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$ and $-X^5S(O)_2R^{13}$ and/or 1 radical selected from $-R^{14}$, $-X^5OR^{14}$, $-X^5SR^{14}$, $-X^5S(O)R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$, $-X^5OC(O)R^{14}$, $-X^5NR^{14}R^{12}$, $-X^5NR^{12}C(O)R^{14}$, $-X^5NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{14}R^{12}$, $-X^5NR^{12}S(O)_2R^{14}$, $-X^5NR^{12}C(O)NR^{14}R^{12}$ and $-X^5NR^{12}C(NR^{12})NR^{14}R^{12}$, wherein X^5 is a bond or (C_{1-6}) alkylene; R^{12} at each occurrence independently is hydrogen, (C_{1-6}) alkyl or halo-substituted (C_{1-6}) alkyl; R^{13} is (C_{1-6}) alkyl or halo-substituted (C_{1-6}) alkyl; and R^{14} is (C_{3-10}) cycloalkyl (C_{0-6}) alkyl, hetero (C_{3-10}) cycloalkyl (C_{0-3}) alkyl, (C_{6-10}) aryl (C_{0-6}) alkyl, hetero (C_{5-10}) aryl (C_{0-6}) alkyl, (C_{9-10}) bicycloaryl (C_{0-6}) alkyl or hetero (C_{8-10}) bicycloaryl (C_{0-6}) alkyl;

R^1 is hydrogen and R^2 is (C_{1-6}) alkyl; and

R^3 is $-CH_2X^6$, wherein X^6 is $-X^5NR^{12}S(O)_2R^{12}$ or $-X^5S(O)_2R^{14}$ wherein X^5 , R^{12} and R^{14} are as defined above;

wherein within R^3 any alicyclic or aromatic ring system is unsubstituted or substituted further by 1 to 5 radicals independently selected from (C_{1-6}) alkyl, (C_{1-6}) alkylidene, cyano, halo, halo-substituted (C_{1-4}) alkyl, nitro, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$, $-X^5NR^{12}C(O)OR^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$, $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$, $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$, $-X^5C(O)R^{13}$ and $-X^5S(O)_2R^{13}$ and within R^3 any aliphatic moiety is unsubstituted or substituted further by 1 to 5 radicals independently selected from cyano, halo, nitro, $-NR^{12}R^{12}$, $-NR^{12}C(O)R^{12}$, $-NR^{12}C(O)OR^{12}$, $-NR^{12}C(O)NR^{12}R^{12}$, $-NR^{12}C(NR^{12})NR^{12}R^{12}$, $-OR^{12}$, $-SR^{12}$, $-C(O)OR^{12}$,

-27-

-C(O)R¹², -OC(O)R¹², -C(O)NR¹²R¹², -S(O)₂NR¹²R¹², -NR¹²S(O)₂R¹², -P(O)(OR¹²)OR¹²,
 -OP(O)(OR¹²)OR¹², -NR¹²C(O)R¹³, -S(O)R¹³ and -S(O)₂R¹³; wherein X⁵, R¹², R¹³ and R¹⁴ are
 as described above; with the proviso that only one bicyclic ring structure is present within R³;
 and the *N*-oxide derivatives, prodrug derivatives, protected derivatives, individual isomers and
 5 mixtures of isomers thereof; and the pharmaceutically acceptable salts and solvates of such
 compounds and the *N*-oxide derivatives, prodrug derivatives, protected derivatives, individual
 isomers and mixtures of isomers thereof.

Preferred are compounds of the invention in which X¹ is -NHC(R¹)(R²)X³ or
 -NHCH(R¹⁹)C(O)R²⁰, wherein R¹ is hydrogen or (C₁₋₆)alkyl and R² is hydrogen, (C₁₋₆)alkyl,
 10 -X⁵OR¹², -X⁵S(O)R¹³, -X⁵OR¹⁴, (C₆₋₁₀)aryl(C₀₋₆)alkyl or hetero(C₅₋₁₀)aryl(C₀₋₆)alkyl or R¹ and
 R² taken together with the carbon atom to which both R¹ and R² are attached form
 (C₃₋₆)cycloalkylene or (C₃₋₆)heterocycloalkylene, wherein within said R² any heteroaryl, aryl,
 cycloalkylene or heterocycloalkylene is unsubstituted or substituted with (C₁₋₆)alkyl or
 hydroxy, particularly wherein X³ is cyano, -C(O)R¹⁶, -C(R⁶)(OR⁶)₂, -CH=CHS(O)₂R⁵,
 15 -CH₂C(O)R¹⁶, -C(O)CF₂C(O)NR⁵R⁵, -C(O)C(O)NR⁵R⁶, -C(O)C(O)OR⁵, -C(O)CH₂OR⁵,
 -C(O)CH₂N(R⁶)SO₂R⁵ or -C(O)C(O)R⁵, wherein R⁵, R⁶ and R¹⁶ are as described above, and
 R¹⁹ and R²⁰ together with the atoms to which R¹⁹ and R²⁰ are attached form
 (C₄₋₈)heterocycloalkylene, wherein no more than one of the ring member atoms comprising
 the ring is a heteroatom selected from -NR²¹- or -O-, particularly wherein the ring is
 20 unsubstituted or substituted with (C₁₋₆)alkyl or -X⁵C(O)OR¹² and R²¹ is hydrogen, (C₁₋₆)alkyl,
 -X⁵C(O)R¹², -X⁵C(O)OR¹², -R¹⁴, -X⁵C(O)R¹⁴ or -C(O)OR¹⁴.

Particularly preferred are compounds of the invention in which X³ is cyano, -C(O)X⁴,
 -C(O)H, -C(O)N(CH₃)OCH₃, -CH(OCH₃)₂, -C(O)CF₃, -C(O)CF₂CF₃, -CH₂C(O)R¹⁶,
 (E)-2-benzenesulfonyl-vinyl, 2-dimethylcarbamoyl-2,2-difluoro-acetyl, 2-oxo-2-pyrrolidin-1-
 25 yl-acetyl, 2-morpholin-4-yl-2-oxo-acetyl, 2-oxo-2-piperazin-1-yl-acetyl, 2-(4-
 methanesulfonyl-piperazin-1-yl)-2-oxo-acetyl, 2-(1,1-dioxo-1λ⁶-thiomorpholin-4-yl)-2-oxo-
 acetyl, dimethylaminooxalyl, tetrahydro-pyran-4-ylaminooxalyl, 2-morpholin-4-yl-
 ethylaminooxalyl, cyclopentyl-ethyl-aminooxalyl, pyridin-3-ylaminooxalyl,
 phenylaminooxalyl, 1-benzoyl-piperidin-4-ylaminooxalyl, 1-benzylcarbamoyl-methanoyl,
 30 1-benzyloxy(oxalyl), 2-benzyloxy-acetyl, 2-benzenesulfonylamino-ethanoyl,
 2-oxo-2-phenyl-ethanoyl, 3*H*-oxazole-2-carbonyl, 5-trifluoromethyl-oxazole-2-carbonyl, 3-
 trifluoromethyl-[1,2,4]oxadiazole-5-carbonyl, 2,2,3,3,3-pentafluoro-propionyl,
 hydroxyaminooxalyl, oxalyl, 2-(1,3-dihydro-isoindol-2-yl)-2-oxo-acetyl, benzothiazol-2-

ylaminooxalyl, 2-oxo-ethyl, 2-oxazol-2-yl-2-oxo-ethyl or 2-benzooxazol-2-yl-2-oxo-ethyl, particularly wherein X^4 is 1*H*-benzimidazol-2-yl, pyrimidin-2-yl, benzooxazol-2-yl, benzothiazol-2-yl, pyridazin-3-yl, 3-phenyl-[1,2,4]oxadiazol-5-yl, 5-ethyl-[1,3,4]-oxadiazol-2-yl, 5-ethyl-[1,2,4]-oxadiazol-3-yl or 3-ethyl-[1,2,4]oxadiazol-5-yl; and R^{19} and R^{20} together with the atoms to which R^{19} and R^{20} are attached form 1-benzoyl-3-oxo-piperidin-4-yl, 1-benzoyl-3-oxo-azepan-4-yl, 2-methyl-4-oxo-tetrahydro-furan-3-yl, 2-ethyl-4-oxo-tetrahydro-furan-3-yl, 4-oxo-1-(1-phenyl-methanoyl)-pyrrolidin-3-yl or (S)-2-acetoxy-4-oxo-azetidin-3-yl.

Most particularly preferred are compounds of the invention in which X^3 is $-C(O)X^4$, in particular 1*H*-benzimidazol-2-ylcarbonyl, pyrimidin-2-ylcarbonyl, benzooxazol-2-ylcarbonyl, benzothiazol-2-ylcarbonyl, pyridazin-3-ylcarbonyl, 3-phenyl-[1,2,4]oxadiazol-5-ylcarbonyl, 5-ethyl-[1,2,4]-oxadiazol-3-ylcarbonyl, 5-ethyl-[1,3,4]-oxadiazol-2-ylcarbonyl or 3-ethyl-[1,2,4]oxadiazol-5-ylcarbonyl, or $-C(O)C(O)NR^5R^6$, in particular 2-oxo-2-pyrrolidin-1-yl-acetyl, 2-morpholin-4-yl-2-oxo-acetyl, 2-oxo-2-piperazin-1-yl-acetyl, 2-(4-methanesulfonyl-piperazin-1-yl)-2-oxo-acetyl, 2-(1,1-dioxo-1 λ^6 -thiomorpholin-4-yl)-2-oxo-acetyl, dimethylaminooxalyl, tetrahydro-pyran-4-ylaminooxalyl, 2-morpholin-4-yl-ethylaminooxalyl, cyclopentyl-ethyl-aminooxalyl, pyridin-3-ylaminooxalyl, phenylaminooxalyl or 1-benzoyl-piperidin-4-ylaminooxalyl. Preferred are compounds of the invention in which X^2 is $-OH$ or $-OC(O)NR^{12}R^{12}$, particularly wherein each R^{12} independently represent hydrogen or (C_{1-6}) alkyl, wherein said alkyl is unsubstituted or substituted with hydroxy or methoxy, or X^2 is $-OC(O)NHR^{14}$, wherein R^{14} is (C_{3-10}) cycloalkyl (C_{0-6}) alkyl or hetero (C_{3-10}) cycloalkyl (C_{1-3}) alkyl, or X^2 is $-OC(O)R^{14}$, wherein R^{14} is $-NR^{22}R^{23}$ and R^{22} and R^{23} together with the nitrogen atom to which both R^{22} and R^{23} attached form a hetero (C_{4-6}) cycloalkyl ring, which ring may be unsubstituted or substituted with hydroxy, particularly in which X^2 is selected from $-OH$, dimethylcarbamoxyloxy, morpholin-4-ylcarbonyloxy, piperidin-1-yl-carbonyloxy, pyrrolidin-1-yl-carbonyloxy, pyrimidin-2-ylamino, tetrahydro-pyran-4-ylamino, 1-methyl-piperidin-4-ylamino, *N*-(2-methoxyethyl)-*N*-(tetrahydro-pyran-4-yl)amino, isopropylamino and cyclohexylamino, 4-*tert*-butoxycarbonylpiperazin-1-ylcarbonyloxy, *N*-benzyl-carbamoyloxy, pyrrolidin-1-yl-carbonyloxy, *N,N*-dimethyl-carbamoyloxy, piperidin-1-yl-carbonyloxy, 4-methanesulfonyl-piperazin-1-yl-carbonyloxy, 4-ethoxycarbonylpiperazin-1-ylcarbonyloxy, *N*-cyclohexyl-carbamoyloxy, *N*-phenyl-carbamoyloxy, *N*-(5,6,7,8-tetrahydro-naphthalen-1-yl)-carbamoyloxy, *N*-butyl-*N*-methyl-carbamoyloxy, *N*-pyridin-3-yl-carbamoyloxy, *N*-isopropyl-carbamoyloxy, *N*-pyridin-4-yl-carbamoyloxy, *N*-cyanomethyl-*N*-methyl-carbamoyloxy, *N,N*-bis-(2-methoxy-ethyl)-carbamoyloxy, *N*-phenethyl-carbamoyloxy, piperazine- carbonyloxy,

N-naphthalen-2-yl-carbamoyloxy, 4-benzyl-piperazine-1-carbamoyloxy, 4-(1-furan-2-yl-carbonyl)-piperazine-1-carbamoyloxy, thiomorpholin-4-yl-carbonyloxy, 1,1-dioxo-1 λ ⁶-thiomorpholin-4-yl-carbonyloxy, bis-(2-methoxy-ethyl)-carbamoyloxy, morpholin-4-ylcarbonyloxy, 2-methoxyethylcarbamoyloxy, diethylcarbamoyloxy, pyrrolidin-5
1-ylcarbonyloxy, 2-hydroxyethylcarbamoyloxy, tetrahydro-furan-2-ylmethylcarbamoyloxy, cyclopropylcarbamoyloxy, *tert*-butylcarbamoyloxy, 3-hydroxy-pyrrolidin-1-yl-carbonyloxy and carbamoyloxy, more particularly morpholin-4-ylcarbonyloxy, 2-methoxyethylcarbamoyloxy, diethylcarbamoyloxy, pyrrolidin-1-ylcarbonyloxy, 2-hydroxyethylcarbamoyloxy, tetrahydro-furan-2-ylmethylcarbamoyloxy, cyclopropylcarbamoyloxy, *tert*-butylcarbamoyloxy, 3-hydroxy-pyrrolidin-1-yl-carbonyloxy and carbamoyloxy.

Preferred are compounds of the invention in which X² is -NHR¹⁵, wherein R¹⁵ is (C₆₋₁₀)aryl, hetero(C₅₋₁₀)aryl, (C₉₋₁₀)bicycloaryl or hetero(C₈₋₁₀)bicycloaryl, or -NR¹⁷R¹⁸, wherein R¹⁷ is hetero(C₃₋₁₀)cycloalkyl and R¹⁸ is hydrogen or R¹⁷ and R¹⁸ independently are 15 (C₆₋₁₀)aryl(C₁₋₆)alkyl or hetero(C₅₋₁₀)aryl(C₁₋₆)alkyl, wherein within R¹⁵, R¹⁷ and R¹⁸ any alicyclic or aromatic ring system is unsubstituted or substituted further by 1 to 5 radicals independently selected from (C₁₋₆)alkyl, cyano, halo, nitro, halo-substituted(C₁₋₄)alkyl, -X⁵OR¹², -X⁵C(O)OR¹², -X⁵C(O)R¹³, -X⁵C(O)NR¹²R¹², -X⁵NR¹²S(O)₂R¹² and/or 1 radical selected from -R¹⁴, -X⁵OR¹⁴ and -X⁵C(O)NR¹⁴R¹², in particular in which X² is selected from 20 5-nitrothiazol-2-ylamino, 2-nitrophenylamino, pyrimidin-2-ylamino, tetrahydro-pyran-4-ylamino, *N*-(2-methoxyethyl)-*N*-(tetrahydro-pyran-4-yl)amino, 1-methyl-piperidin-4-ylamino, isopropylamino, di(thien-2-ylmethyl)amino or di(benzyl)amino.

Preferred are compounds of the invention in which X² is -OR⁴ wherein R⁴ is 4-methoxy-phenyl, 4'-hydroxymethyl-phenyl, methoxymethyl, phenyl-methanoyl, 1-(4-25 phenoxy-phenyl)-methanoyl, 3-biphenyl, 4-biphenyl, 1-biphenyl-4-yl-methanoyl, naphthalen-2-yl-methanoyl, benzo[1,3]dioxol-5-yl-methanoyl, (4-methanesulfonylamino-phenyl)-methanoyl, benzo[*b*]thien-2-yl-methanoyl, 4'-chloro-4-biphenyl, 4-hydroxy-phenyl-methanoyl, 3-chloro-benzo[*b*]thien-2-yl-methanoyl, thien-2-yl-methanoyl, thien-3-yl-methanoyl, 3-chloro-thien-2-yl-methanoyl, 5-methyl-thien-2-yl-methanoyl, 4-methoxy-phenyl 30 methanoyl, 4-trifluoromethoxy-phenyl methanoyl, 4-chloro-phenyl-methanoyl, 3-bromo-phenyl, cyclohexylmethyl, 3,4-dimethoxy-phenyl-methanoyl, 3,4-difluorophenyl-methanoyl, 3-fluoro, 4-methoxy-phenyl-methanoyl, 4-fluorophenyl-methanoyl, 4-trifluoromethyl-phenyl-methanoyl, 4-formyl-phenyl-formyl, 3-formyl-phenyl-formyl, 4-methyl-pentanoyl, tetrahydro-

pyran-4-ylmethyl 2-morpholin-4-yl-2-oxo-ethyl.

Most particularly preferred are compounds of the invention in which X^2 is selected from -OH, dimethylcarbamoyloxy, morpholin-4-ylcarbonyloxy, piperidin-1-yl-carbonyloxy, pyrrolidin-1-yl-carbonyloxy, pyrimidin-2-ylamino, tetrahydro-pyran-4-ylamino, 1-methyl-
 5 piperidin-4-ylamino, *N*-(2-methoxyethyl)-*N*-(tetrahydro-pyran-4-yl)amino, isopropylamino and cyclohexylamino.

Preferred are compounds of the invention in which R^1 is hydrogen or (C_{1-6}) alkyl and R^2 is hydrogen, $-X^5OR^{12}$, $-X^5R^{12}$, (C_{5-10}) heteroaryl (C_{0-6}) alkyl, (C_{5-10}) aryl (C_{0-6}) alkyl, (C_{5-10}) cycloalkyl (C_{0-6}) alkyl, (C_{5-10}) heterocycloalkyl (C_{0-6}) alkyl or (C_{1-6}) alkyl; or R^1 and R^2
 10 taken together with the carbon atom to which both R^1 and R^2 are attached form (C_{3-8}) cycloalkylene or (C_{3-8}) heterocycloalkylene; wherein within said R^2 any heteroaryl, aryl, cycloalkyl, heterocycloalkyl, cycloalkylene or heterocycloalkylene is optionally substituted with 1 to 3 radicals independently selected from (C_{1-6}) alkyl and hydroxy; particularly in which R^1 is hydrogen or methyl and R^2 is hydrogen, methoxymethyl, (C_{1-6}) alkyl, phenethyl,
 15 thien-2-yl or 5-methyl-furan-2-yl or R^1 and R^2 taken together with the carbon atom to which both R^1 and R^2 are attached form cyclopropylene, tetrahydro-pyran-4-ylene or methyl-piperidin-4-ylene.

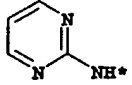
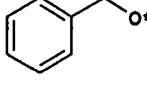

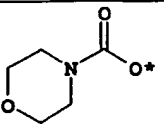
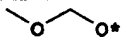
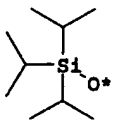
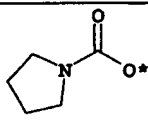
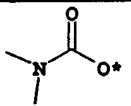
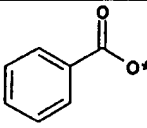
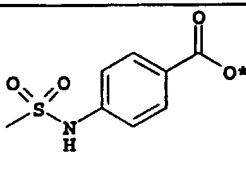
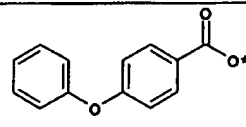
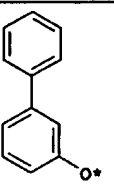
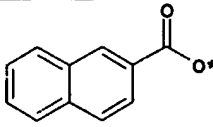
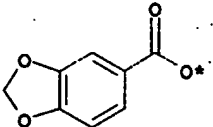
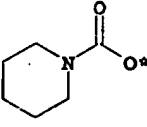
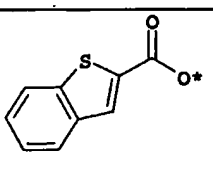
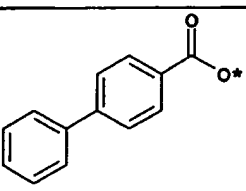
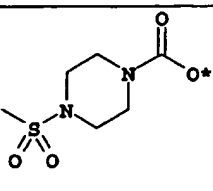
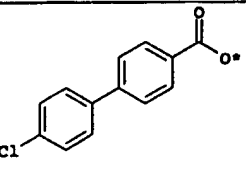
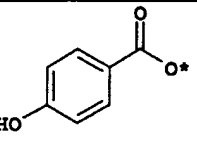
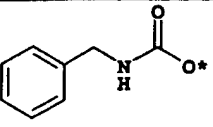
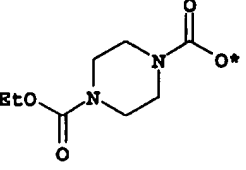
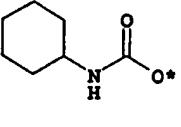
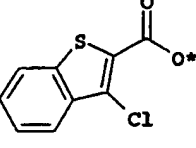
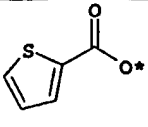
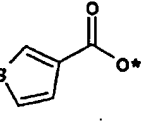
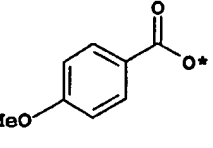
Preferred are compounds of the invention in which R^3 is $-CH_2X^6$; wherein X^6 is selected from $-X^5SR^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2R^{13}$, $-X^5C(O)R^{13}$, $-X^5OR^{12}$, $-X^5SR^{14}$, $-X^5R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$,
 20 $-X^5C(O)NR^{14}R^{12}$, wherein X^5 , R^{12} , R^{13} and R^{14} are as defined above; particularly wherein R^3 is thiophene-2-sulfonyl-methyl, 3-chloro-2-fluoro-phenyl-methane-sulfonyl-methyl, benzene-sulfonyl-methyl, phenyl-methane-sulfonyl-methyl, 2-(1,1-difluoro-methoxy)-phenyl-methane-sulfonyl-methyl, 2-benzene-sulfonyl-ethyl, 2-(pyridine-2-sulfonyl)-ethyl, 2-(pyridine-4-sulfonyl)-ethyl, 2-phenyl-methanesulfonyl-ethyl, oxy-pyridin-2-yl-methane-sulfonyl-methyl, prop-2-ene-1-sulfonyl-methyl, 4-methoxy-phenyl-methane-sulfonyl-
 25 methyl, *p*-tolyl-methane-sulfonyl-methyl, 4-chloro-phenyl-methane-sulfonyl-methyl, *o*-tolyl-methane-sulfonyl-methyl, 3,5-dimethyl-phenyl-methane-sulfonyl-methyl, 4-trifluoro-methyl-phenyl-methane-sulfonyl-methyl, 4-trifluoro-methoxy-phenyl-methane-sulfonyl-methyl, 2-bromo-phenyl-methane-sulfonyl-methyl, pyridin-2-yl-methane-sulfonyl-methyl, pyridin-3-yl-methane-sulfonyl-methyl, pyridin-4-yl-methane-sulfonyl-methyl, naphthalen-2-yl-methane-sulfonyl-methyl, 3-methyl-phenyl-methane-sulfonyl-methyl, 3-trifluoro-
 30 methyl-phenyl-methane-sulfonyl-methyl, 3-trifluoro-methoxy-phenyl-methane-sulfonyl-methyl, 4-fluoro-2-trifluoromethoxy-phenyl-methane-sulfonylmethyl, 2-fluoro-6-trifluoromethyl-phenylmethanesulfonylmethyl, 3-chloro-phenylmethanesulfonylmethyl, 2-fluoro-phenylmethanesulfonylmethyl, 2-trifluoro-phenylmethanesulfonylmethyl, 2-cyano-phenylmethanesulfonylmethyl, 4-*tert*-butyl-phenylmethanesulfonylmethyl, 2-fluoro-3-methyl-phenyl-
 35 methane-sulfonyl-methyl, 3-fluoro-phenylmethanesulfonylmethyl, 4-fluoro-phenylmethane-sulfonylmethyl, 2-chloro-phenylmethanesulfonylmethyl, 2,5-difluoro-phenylmethane-sulfonylmethyl,

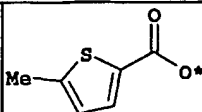
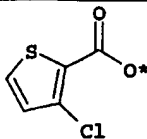
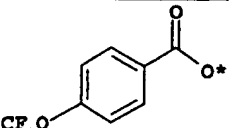
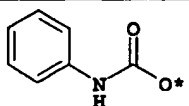
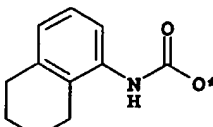
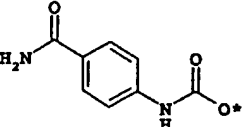
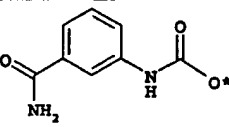
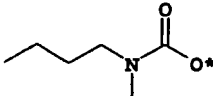
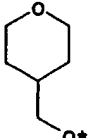
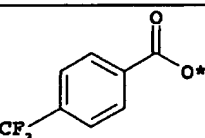
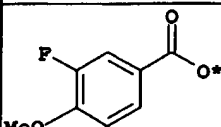
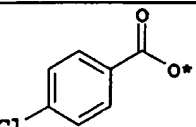
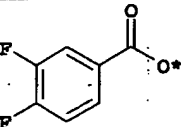
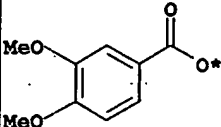
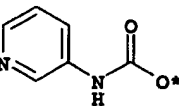
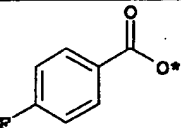

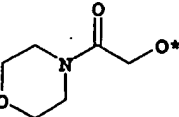
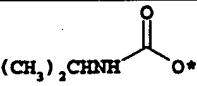
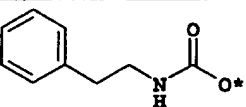
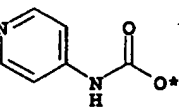
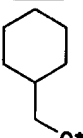
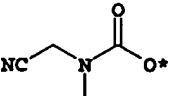
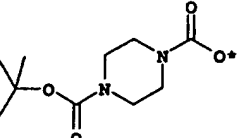
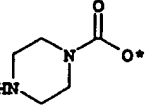
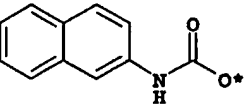
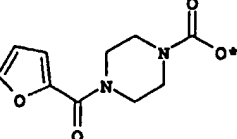
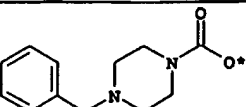
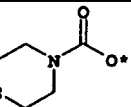
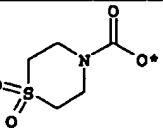
2,6-difluoro-phenylmethanesulfonylmethyl, 2,5-dichloro-phenyl-methane-sulfonylmethyl,
 3,4-dichloro-phenylmethanesulfonylmethyl, 2-(1,1-difluoro-methoxy)-phenyl-methanesulfonylmethyl,
 2-cyano-phenyl-methane-sulfonyl-methyl, 3-cyano-phenylmethanesulfonylmethyl, 2-trifluoro-methoxy-phenyl-
 methane-sulfonylmethyl, 2,3-difluoro-phenylmethanesulfonylmethyl, 2,5-difluoro-phenyl-
 5 methanesulfonylmethyl, biphenyl-2-ylmethanesulfonylmethyl, cyclohexylmethyl, 3-fluoro-phenyl-
 methanesulfonylmethyl, 3,4-difluoro-phenyl-methanesulfonylmethyl,
 2,4-difluoro-phenylmethanesulfonylmethyl, 2,4,6-trifluoro-phenylmethanesulfonylmethyl,
 2,4,5-trifluoro-phenylmethanesulfonylmethyl, 2,3,4-trifluoro-phenylmethanesulfonylmethyl,
 2,3,5-trifluoro-phenyl-methane-sulfonylmethyl, 2,5,6-trifluoro-phenylmethanesulfonylmethyl,
 10 2-chloro-5-trifluoro-methylphenylmethanesulfonylmethyl, 2-methyl-propane-1-sulfonyl, 2-fluoro-3-trifluoro-
 methylphenylmethanesulfonylmethyl, 2-fluoro-4-trifluoro-methylphenylmethanesulfonylmethyl,
 2-fluoro-5-trifluoro-methyl-phenyl-methane-sulfonyl-methyl, 4-fluoro-3-trifluoro-
 methylphenylmethanesulfonylmethyl, 2-methoxy-phenyl-methanesulfonylmethyl,
 3,5-bis-trifluoromethyl-phenylmethanesulfonylmethyl, 4-difluoromethoxy-phenylmethanesulfonylmethyl,
 15 2-difluoro-methoxy-phenyl-methanesulfonylmethyl, 3-difluoromethoxy-phenylmethanesulfonylmethyl,
 2,6-dichloro-phenylmethanesulfonylmethyl, biphenyl-4-ylmethanesulfonylmethyl,
 3,5-dimethyl-isoxazol-4-ylmethanesulfonylmethyl, 5-chloro-thien-2-yl-methane-sulfonylmethyl,
 2-[4-(1,1-difluoro-methoxy)-benzenesulfonyl]-ethyl, 2-[2-(1,1-difluoro-methoxy)-benzenesulfonyl]-ethyl,
 2-[3-(1,1-difluoro-methoxy)-benzenesulfonyl]-ethyl, 2-(4-trifluoromethoxy-benzenesulfonyl)-ethyl,
 20 2-(3-trifluoromethoxy-benzenesulfonyl)-ethyl, 2-(2-trifluoro-methoxy-benzene-sulfonyl)-ethyl,
 (cyanomethyl-methyl-carbamoyl)-methyl, biphenyl-3-ylmethyl, 2-oxo-2-pyrrolidin-1-yl-ethyl,
 2-benzenesulfonyl-ethyl, isobutylsulfanylmethyl, 2-phenylsulfanyl-ethyl, cyclohexylmethanesulfonylmethyl,
 2-cyclohexyl-ethanesulfonyl, benzyl, naphthalen-2-yl, benzylsulfanylmethyl,
 2-trifluoromethyl-benzylsulfanylmethyl, phenylsulfanyl-ethyl, cyclopropyl-methanesulfonylmethyl, 5-bromo-
 25 thien-2-ylmethyl, 3-phenyl-propyl, 2,2-difluoro-3-phenyl-propyl, 3,4,5-trimethoxy-
 phenylmethanesulfonylmethyl, 2,2-difluoro-3-thien-2-yl-propyl, cyclohexylethyl, cyclohexylmethyl, *tert*-
 butylmethyl, 1-methylcyclohexylmethyl, 1-methylcyclopentylmethyl, 2,2-difluoro-3-phenylpropyl, 2,2-dimethyl-
 3-phenylpropyl, 1-benzylcyclopropylmethyl, $-X^5S(O)_2R^{13}$ and $-X^5S(O)_2R^{14}$, wherein R^{13} is alkyl and R^{14} is
 phenyl which phenyl is unsubstituted or substituted.

30 Preferred are compounds of the invention in which R^3 is cyclohexylethyl, cyclohexylmethyl, *tert*-
 butylmethyl, 1-methylcyclohexylmethyl, 1-methylcyclopentylmethyl, 2,2-difluoro-3-phenylpropyl, 2,2-dimethyl-
 3-phenylpropyl, 1-benzylcyclopropylmethyl, $-X^5S(O)_2R^{13}$ or $-X^5S(O)_2R^{14}$, wherein R^{13} is alkyl and R^{14} is phenyl
 which phenyl is unsubstituted or substituted.

35 The following tables are intended to provide guidance to better carry out the present invention.
 However, they do not limit the scope of the invention. People of ordinary skill may selectively make particular
 compounds by joining O*, HN* or H* of one of the fragments (A1 to A62) shown in Table 1 to the methine
 carbon atom (*CH*) of one of the fragments (B1 to B93) shown in Table 2, and joining the methine carbon atom
 (*CH* or *CF*) of one of the fragments (B1 to B93) shown in Table 2 to the acyl carbon atom (C*) of one of the
 40 fragments (C1 to C91) depicted in Table 3.

TABLE 1

A1		A2		A3	
A4		A5		A6	
A7		A8		A9	
A10		A11		A12	
A13		A14		A15	
A16		A17		A18	
A19		A20		A21	
A22		A23		A24	
A25		A26		A27	

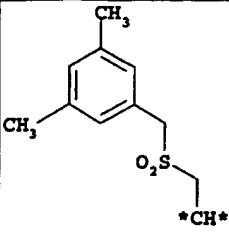
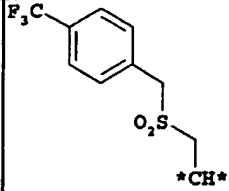
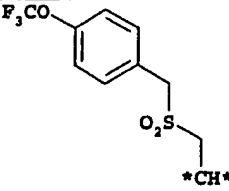
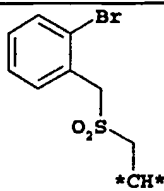
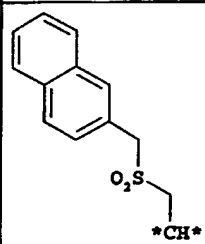
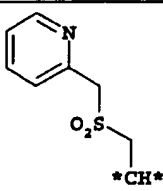
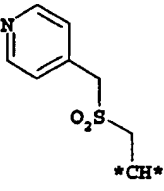
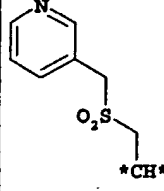
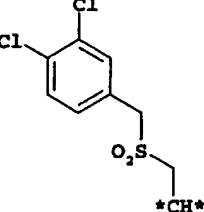
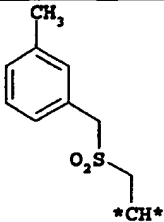
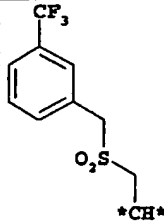
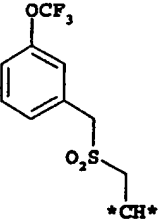
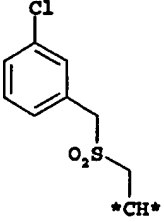
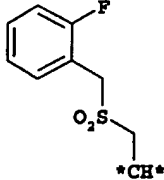
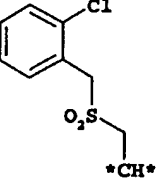
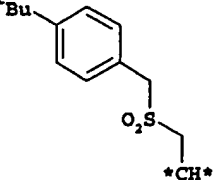
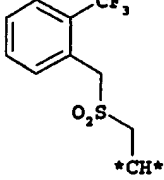
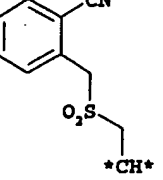
A28		A29		A30	
A31		A32		A33	
A34		A35		A36	
A37		A38		A39	
A40		A41		A42	
A43		A44		A45	
A46		A47		A48	
A49		A50		A51	
A52		A53		A54	
A55		A56		A57	

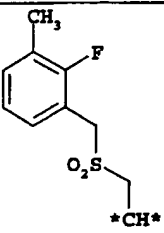
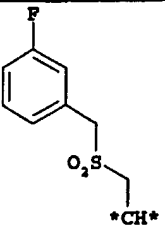
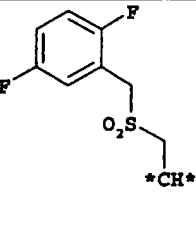
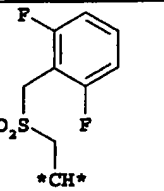
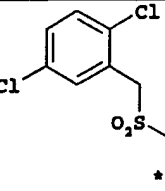
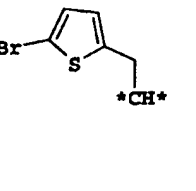
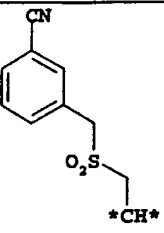
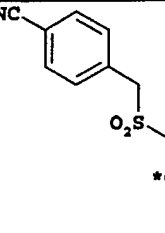
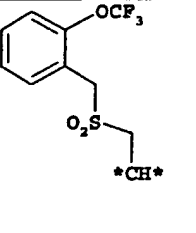
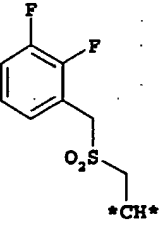
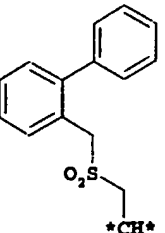
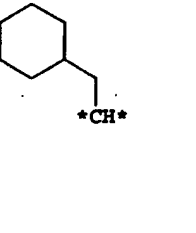
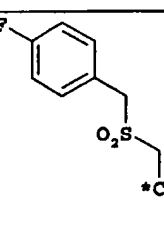
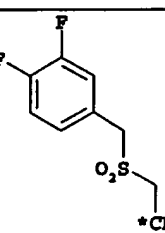
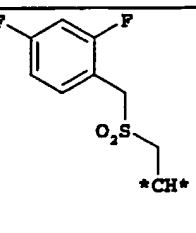
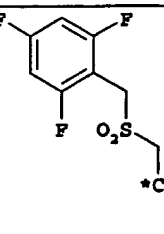
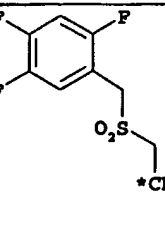
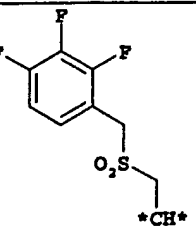
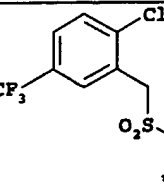
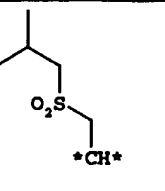
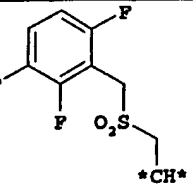
-34-

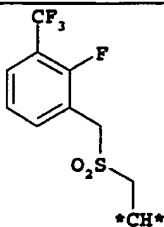
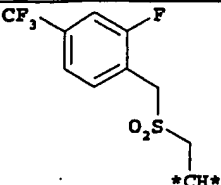
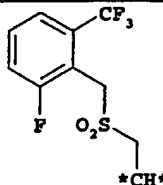
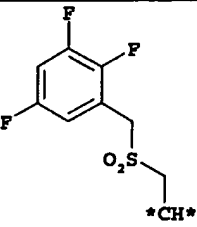
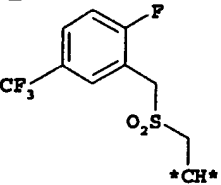
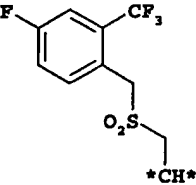
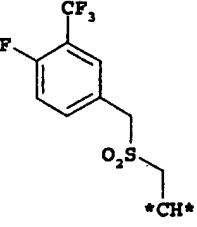
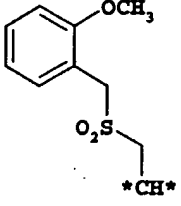
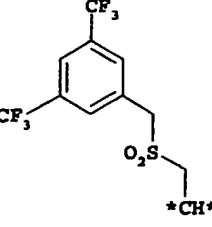
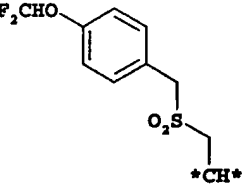
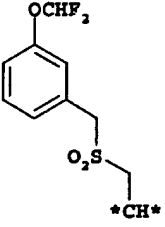
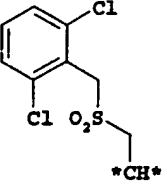
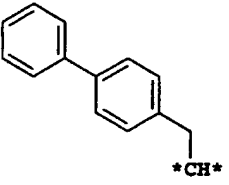
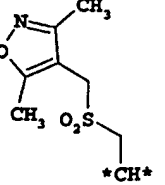
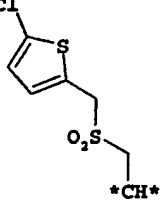
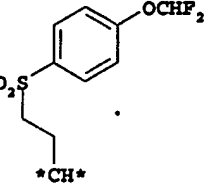
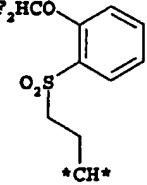
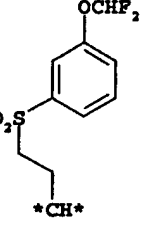
A58		A59		A60	
A61	H*	A62	F*		

TABLE 2

B1		B2		B3	
B4		B5		B6	
B7		B8		B9	
B10		B11		B12	
B13		B14		B15	

B16		B17		B18	
B19		B20		B21	
B22		B23		B24	
B25		B26		B27	
B28		B29		B30	
B31		B32		B33	

B34		B35		B36	
B37		B38		B39	
B40		B41		B42	
B43		B44		B45	
B46		B47		B48	
B49		B50		B51	
B52		B53		B54	

B55		B56		B57	
B58		B59		B60	
B61		B62		B63	
B64		B65		B66	
B67		B68		B69	
B70		B71		B72	

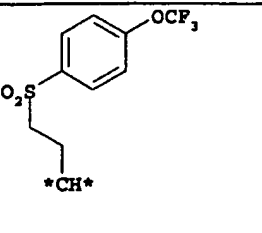
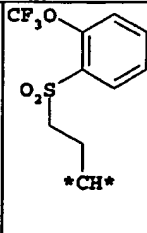
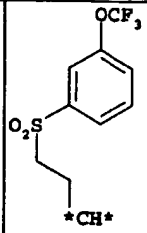
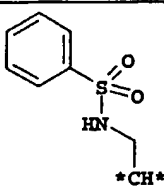
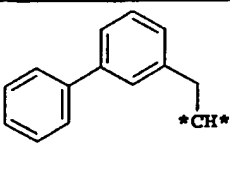
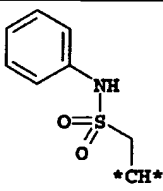
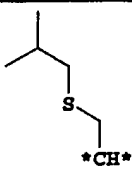
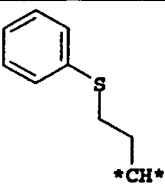
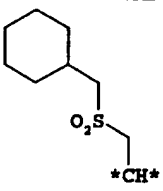
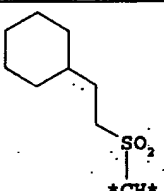
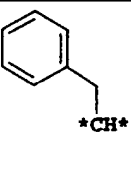
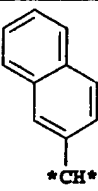
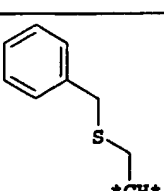
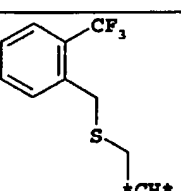
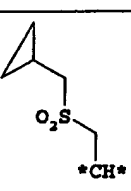
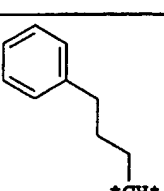
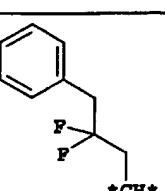
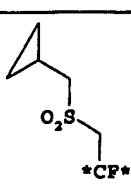
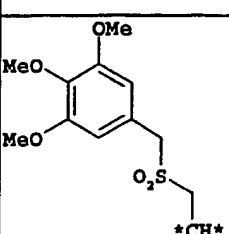
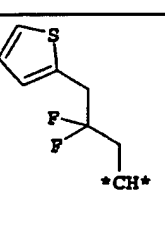
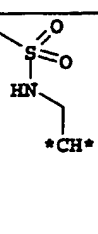
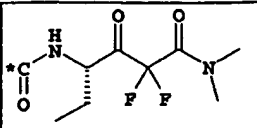
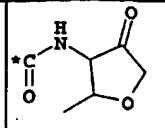
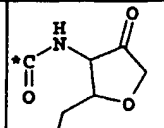
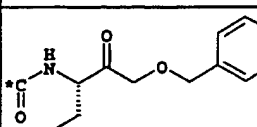
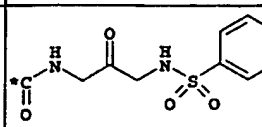
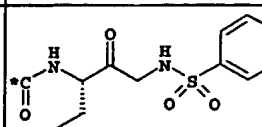
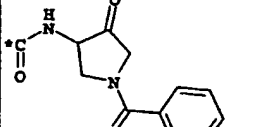
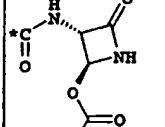
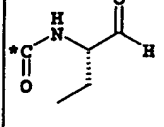
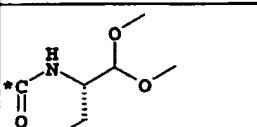
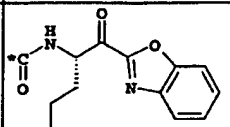
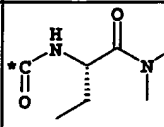
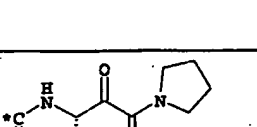
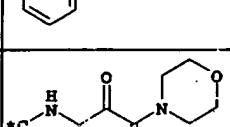
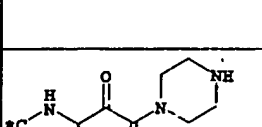
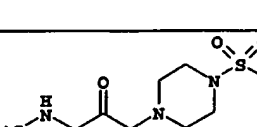
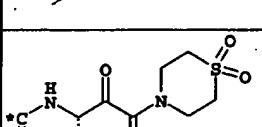
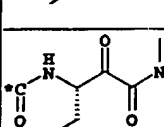
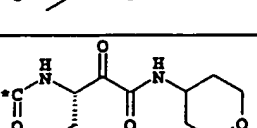
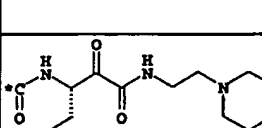
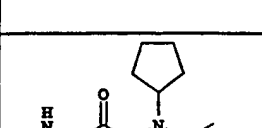
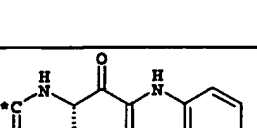
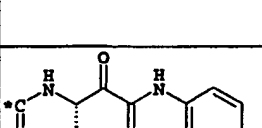
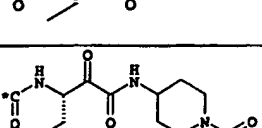
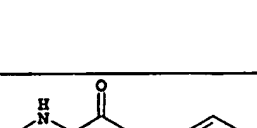
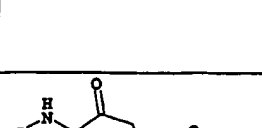
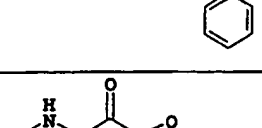
B73		B74		B75	
B76		B77		B78	
B79		B80		B81	
B82		B83		B84	
B85		B86		B87	
B88		B89		B90	
B91		B92		B93	

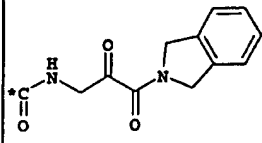
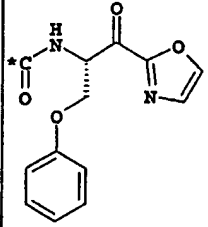
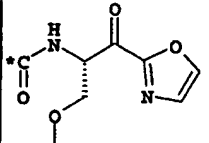
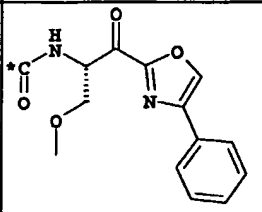
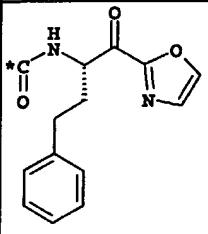
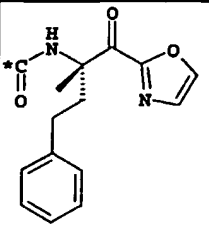
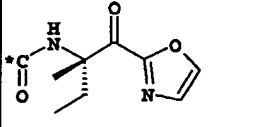
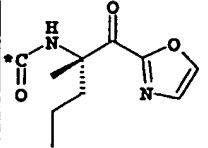
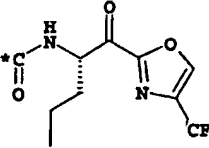
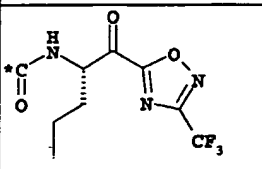
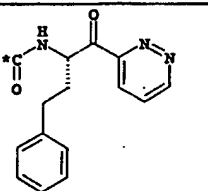
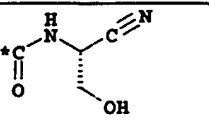
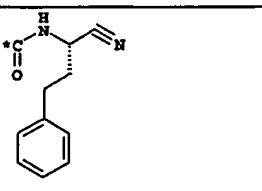
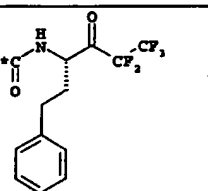
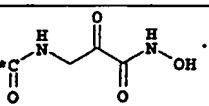
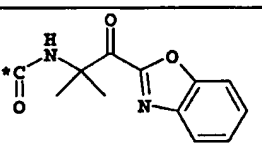
TABLE 3

C1		C2		C3	
C4		C5		C6	
C7		C8		C9	
C10		C11		C12	
C13		C14		C15	
C16		C17		C18	
C19		C20		C21	
C22		C23		C24	

C25		C26		C27	
C28		C29		C30	
C31		C32		C33	
C34		C35		C36	
C37		C38		C39	
C40		C41		C42	
C43		C44		C45	
C46		C47		C48	
C49		C50		C51	

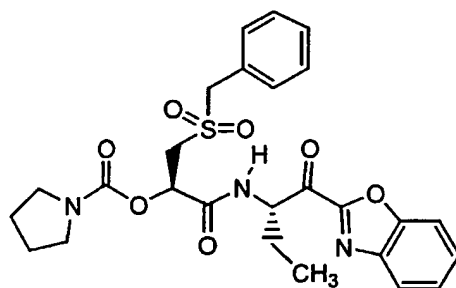
C52		C53		C54	
C55		C56		C57	
C58		C59		C60	
C61		C62		C63	
C64		C65		C66	
C67		C68		C69	
C70		C71		C72	
C73		C74		C75	

-42-

C76		C77		C78	
C79		C80		C81	
C82		C83		C84	
C85		C86		C87	
C88		C89		C90	
C91					

For convenience, compounds of the present invention may be referenced to by their "A", "B", and "C" fragment combinations. Thus, for example, the compound referenced as A7-B4-C13 is the product of the combination of group A7 in Table 1 and B4 in Table 2 and C13 in Table 3, namely pyrrolidine-1-carboxylic acid (R)-1-[(S)-1-(1-benzooxazol-2-yl-methanoyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester:

-43-



Further preferred compounds of Formula I are provided in the following:

- (*R*)-*N*-cyanomethyl-2-hydroxy-3-phenylmethanesulfonyl-propionamide;
- 5 (*R*)-*N*-(1-cyano-1-thiophen-2-yl-methyl)-2-hydroxy-3-phenylmethanesulfonyl-propionamide;
- (*R*)-*N*-(1-cyano-1-thiophen-2-yl-methyl)-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionamide;
- (*R*)-*N*-cyanomethyl-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionamide;
- 10 morpholine-4-carboxylic acid (*R*)-1-(cyanomethyl-carbamoyl)-2-phenylmethanesulfonyl-ethyl ester;
- morpholine-4-carboxylic acid (*R*)-1-(cyanomethyl-carbamoyl)-2-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-ethyl ester;
- (*R*)-(2-methoxy-ethyl)-carbamic acid 1-(cyanomethyl-carbamoyl)-2-phenylmethanesulfonyl-ethyl ester;
- 15 (*S*)-diethyl-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (*S*)-pyrrolidine-1-carboxylic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (*S*)-morpholine-4-carboxylic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (*S*)-4-Ethyl-piperazine-1-carboxylic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (*S*)-2-hydroxymethyl-pyrrolidine-1-carboxylic acid (*S*)-1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- 20 (*S*)-(2,2,2-Trifluoro-ethyl)-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (*S*)-(2-hydroxyethyl)-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (Tetrahydrofuran-2-ylmethyl)-carbamic acid (*S*)-1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (*S*)-Azetidine-1-carboxylic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- 25 (*S*)-cyclopropyl-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (*S*)-piperidine-1-carboxylic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (*S*)-(2-methoxy-ethyl)-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (*R*)-3-hydroxy-pyrrolidine-1-carboxylic acid (*S*)-1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (*S*)-3-hydroxy-pyrrolidine-1-carboxylic acid (*S*)-1-(cyanomethyl-carbamoyl)-2-cyclohexyl-
- 30 ethyl ester;

- (S)-morpholine-4-carboxylic acid 1-(cyanomethyl-carbamoyl)-3-cyclohexyl-propyl ester;
 morpholine-4-carboxylic acid (*R*)-1-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-
 propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester;
 morpholine-4-carboxylic acid (*R*)-1-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-
 5 propylcarbamoyl]-2-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-ethyl ester;
 morpholine-4-carboxylic acid (*R*)-1-[(*S*)-1-(1-benzothiazol-2-yl-methanoyl)-
 propylcarbamoyl]-2-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-ethyl ester;
 pyrrolidine-1-carboxylic acid (*R*)-1-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-
 propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester;
 10 dimethyl-carbamic acid (*R*)-1-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propylcarbamoyl]-2-
 phenylmethanesulfonyl-ethyl ester;
 morpholine-4-carboxylic acid (*R*)-1-[(*S*)-1-(1-benzylcarbamoyl-methanoyl)-
 propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester;
 morpholine-4-carboxylic acid (*S*)-1-[(*S*)-1-(oxazol[4,5-*b*]pyridine-2-carbonyl)-propylcarbamoyl]-2-
 15 phenylmethanesulfonyl-ethyl ester;
 morpholine-4-carboxylic acid (*S*)-1-[(*S*)-1-(5-ethyl-[1,3,4]oxadiazole-2-carbonyl)-propylcarbamoyl]-2-
 phenylmethanesulfonyl-ethyl ester;
 (*S*)-2-[(*R*)-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-
 propanoylamino]-*N*-methoxy-*N*-methyl-butylamide;
 20 (*R*)-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-*N*-[(*S*)-1-formyl-propyl]-2-hydroxy-
 propionamide;
 (*R*)-*N*-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propyl]-2-hydroxy-3-phenyl-methanesulfonyl-
 propionamide;
 (*S*)-3-{3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-propanoylamino}-2-oxo-
 25 pentanoic acid benzylamide;
N-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propyl]-3-[2-(1,1-difluoro-methoxy)-
 phenylmethanesulfonyl]-propionamide;
N-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-3-phenyl-propyl]-3-*p*-tolylmethanesulfonyl-propionamide;
 3-(2-difluoromethoxy-phenylmethanesulfonyl)-*N*-(1-ethyl-2,3-dioxo-3-pyrrolidin-1-yl-
 30 propyl)-propionamide;
 3-(2-difluoromethoxy-phenylmethanesulfonyl)-*N*-(1-ethyl-3-morpholin-4-yl-2,3-dioxo-
 propyl)-propionamide;
 3-(2-difluoromethoxy-phenylmethanesulfonyl)-*N*-(1-ethyl-2,3-dioxo-3-piperazin-1-yl-propyl)-
 propionamide;

- 3-(2-difluoromethoxy-phenylmethanesulfonyl)-*N*-[3-(1,1-dioxo-1*H*-thiomorpholin-4-yl)-1-ethyl-2,3-dioxo-propyl]-propionamide;
- 3-(2-difluoromethoxy-phenylmethanesulfonyl)-*N*-[1-ethyl-3-(4-methyl-sulfonyl-piperazin-1-yl)-2,3-dioxo-propyl]-propionamide;
- 5 3-[3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid dimethylamide;
- 3-[3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid cyclopentyl-ethyl-amide;
- 3-[3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid
- 10 phenylamide;
- 3-[3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid pyridin-3-ylamide;
- 3-[3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid (tetrahydro-pyran-4-yl)-amide;
- 15 3-[3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid (1-benzoyl-piperidin-4-yl)-amide;
- 3-[3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid (2-morpholin-4-yl-ethyl)-amide;
- (*R*)-*N*-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propyl]-2-(2-nitro-phenylamino)-3-
- 20 phenylmethanesulfonyl-propionamide;
- N*-[1-(benzooxazole-2-carbonyl)-propyl]-3-phenylmethanesulfonyl-2-(pyrimidin-2-ylamino)-propionamide.
- (*R*)-*N*-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-butyl]-2-(5-nitro-thiazol-2-ylamino)-3-phenylmethanesulfonyl-propionamide;
- 25 (2*S*) (4,4-difluoro-2-hydroxy-5-phenyl-pentanoic acid (1(*S*)-cyano-3-phenyl-propyl)-amide;
- N*-(1(*S*)-cyano-3-phenyl-propyl)-2-(*S*)-(2-morpholin-4-yl-2-oxo-ethoxy)-4-phenyl-butyramide;
- N*-(1(*S*)-cyano-3-phenyl-propyl)-2-(*S*)-fluoro-4-phenyl-butyramide;
- N*-(1(*S*)-cyano-3-phenyl-propyl)-2,2-difluoro-4-phenyl-butyramide;
- N*-(1(*S*)-cyano-3-phenyl-propyl)-2-(*S*)-hydroxy-4-phenyl-butyramide;
- 30 *N*-(1(*S*)-cyano-3-phenyl-propyl)-2-(*R*)-hydroxy-4-phenyl-butyramide;
- N*-(1(*S*)-cyano-3-phenyl-propyl)-2-(*R*)-methoxy-4-phenyl-butyramide;
- 2,2-difluoro-5-phenyl-pentanoic acid (1-cyano-cyclopropyl)-amide;
- N*-(1(*S*)-cyano-3-phenyl-propyl)-4-phenyl-butyramide;

- 2,2-difluoro-5-phenyl-pentanoic acid ((S)-1-cyano-3-phenyl-propyl)-amide;
N-(4-cyano-1-ethyl-piperidin-4-yl)-3-cyclohexyl-propionamide;
N-(4-cyano-1-ethyl-piperidin-4-yl)-3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionamide;
(S)-tert-butyl-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
5 (R)-carbamic acid 1-(cyanomethyl-carbamoyl)-2-(2-difluoromethoxy-phenylmethanesulfonyl)-ethyl ester;
(S)-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
(R)-morpholine-4-carboxylic acid 1-(1-cyano-cyclopropylcarbamoyl)-2-phenylmethanesulfonyl-ethyl ester;
(R)-morpholine-4-carboxylic acid 1-(4-cyano-tetrahydro-pyran-4-ylcarbamoyl)-2-phenylmethanesulfonyl-ethyl
10 ester;
3-cyclohexyl-2-hydroxy-*N*-[1-(oxazolo[4,5-*b*]pyridine-2-carbonyl)-propyl]-propionamide;
(R)-*N*-[1-(benzothiazole-2-carbonyl)-butyl]-2-isopropylamino-3-phenylmethanesulfonyl-propionamide;
(R)-*N*-[1-(benzothiazole-2-carbonyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
15 (R)-*N*-[1-(benzothiazole-2-carbonyl)-butyl]-2-dibenzylamino-3-phenylmethanesulfonyl-propionamide;
(R)-*N*-[1-(benzothiazole-2-carbonyl)-butyl]-2-dimethylamino-3-phenylmethanesulfonyl-propionamide;
(R)-*N*-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
20 (R)-*N*-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-(1-methyl-piperidin-4-ylamino)-3-phenylmethanesulfonyl-propionamide;
(R)-*N*-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-(bis-thiophen-2-ylmethyl-amino)-3-phenylmethanesulfonyl-propionamide;
25 (R)-*N*-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-dibenzylamino-3-phenylmethanesulfonyl-propionamide;
(S)-*N*-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-(tetrahydro-pyran-4-ylamino)-3-thiophen-2-yl-propionamide;
(S)-*N*-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-isopropylamino-3-thiophen-2-yl-propionamide;
30 (R)-*N*-[1-(benzothiazole-2-carbonyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
(R)-*N*-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
35

- (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-isopropylamino-3-phenylmethanesulfonyl-propionamide;
- (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-[(2-methoxy-ethyl)-(tetrahydro-pyran-4-yl)-amino]-3-phenylmethanesulfonyl-propionamide;
- 5 (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-cyclohexylamino-3-phenylmethanesulfonyl-propionamide;
- (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-dimethylamino-3-phenylmethanesulfonyl-propionamide;
- (1S)-N-[1-(benzoxazole-2-carbonyl)-butyl]-2-(S)-fluoro-4-phenyl-butyramide;
- 10 2,2-difluoro-5-phenyl-pentanoic acid [(S)-1-(benzoxazole-2-carbonyl)-butyl]-amide;
- morpholine-4-carboxylic acid (S)-1-[(S)-1-(benzoxazole-2-carbonyl)-propylcarbamoyl]-2-cyclohexyl-ethyl ester;
- morpholine-4-carboxylic acid (S)-2-cyclohexyl-1-[(S)-1-(oxazolo[4,5-b]pyridine-2-carbonyl)-propylcarbamoyl]-ethyl ester;
- 15 morpholine-4-carboxylic acid (S)-2-cyclohexyl-1-[(S)-1-(5-ethyl-[1,3,4]oxadiazole-2-carbonyl)-propylcarbamoyl]-ethyl ester;
- morpholine-4-carboxylic acid (S)-2-cyclohexyl-1-[(S)-1-(5-phenyl-[1,3,4]oxadiazole-2-carbonyl)-propylcarbamoyl]-ethyl ester;
- morpholine-4-carboxylic acid (S)-1-[(S)-1-(benzoxazole-2-carbonyl)-propylcarbamoyl]-3-cyclohexyl-propyl ester;
- 20 4-[4,4-dimethyl-2-(morpholine-4-carbonyloxy)-pentanoylamino]-3-oxo-azepane-1-carboxylic acid benzyl ester;
- (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-3-cyclopropylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- 25 (R)-N-[1-(benzoxazole-2-carbonyl)-butyl]-2-cyclohexylamino-3-cyclopropylmethanesulfonyl-propionamide;
- (R)-N-[1-(benzoxazole-2-carbonyl)-butyl]-2-cycloheptylamino-3-cyclopropylmethanesulfonyl-propionamide;
- (R)-3-phenylmethanesulfonyl-N-[(S)-3-phenyl-1-(thiazole-2-carbonyl)-propyl]-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-3-phenyl-propyl]-3-cyclopropylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- 30 (R)-3-cyclopropylmethanesulfonyl-N-[1-(5-ethyl-1,2,4-oxadiazole-3-carbonyl)-propyl]-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- (R)-3-phenylmethanesulfonyl-N-[1-(3-phenyl-1,2,4-oxadiazole-5-carbonyl)-propyl]-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- 35 (R)-N-[1-(3-cyclopropyl-1,2,4-oxadiazole-5-carbonyl)-propyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- {(R)-1-[1-(benzothiazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;

- {(R)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- {(S)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-thiophen-2-yl-ethyl}-carbamic acid tert-butyl ester;
- 5 {(R)-1-[1-(benzothiazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- {(R)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- {(R)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-cyclopropylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- 10 (R)-1-{1-[hydroxy-(3-phenyl-1,2,4-oxadiazol-5-yl)-methyl]-propylcarbamoyl}-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- {(R)-2-cyclopropylmethanesulfonyl-1-[(S)-1-[(5-ethyl-1,2,4-oxadiazol-3-yl)-hydroxy-methyl]-propylcarbamoyl]-ethyl}-carbamic acid tert-butyl ester;
- 15 {(R)-1-[1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- {(R)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-3-phenyl-propylcarbamoyl]-2-cyclopropylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- {(R)-1-[(S)-1-(hydroxy-thiazol-2-yl-methyl)-3-phenyl-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- 20 {(R)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-cyclopropylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- (R)-1-{1-[hydroxy-(3-phenyl-1,2,4-oxadiazol-5-yl)-methyl]-propylcarbamoyl}-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- 25 {(R)-2-cyclopropylmethanesulfonyl-1-[(S)-1-[(5-ethyl-1,2,4-oxadiazol-3-yl)-hydroxy-methyl]-propylcarbamoyl]-ethyl}-carbamic acid tert-butyl ester;
- {(R)-1-[1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- {(R)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-3-phenyl-propylcarbamoyl]-2-cyclopropylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- 30 {(R)-1-[(S)-1-(hydroxy-thiazol-2-yl-methyl)-3-phenyl-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- (R)-2-phenylmethanesulfonyl-1-[(S)-1-[(3-cyclopropyl-1,2,4-oxadiazol-5-yl)-hydroxy-methyl]-propylcarbamoyl]-ethyl}-carbamic acid tert-butyl ester;
- 35 (R)-N-[1-(Benzoxazole-2-carbonyl)-butyl]-2-[cyclopropylmethyl-(tetrahydro-pyran-4-ylmethyl)-amino]-3-phenylmethanesulfonyl-propionamide;
- (R)-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-2-dibenzylamino-3-phenylmethanesulfonyl-propionamide;

- (R)-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- (R)-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-2-isopropylamino-3-phenylmethanesulfonyl-propionamide;
- (R)-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-2-dimethylamino-3-phenylmethanesulfonyl-propionamide;
- 5 (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-(1-methyl-piperidin-4-ylamino)-3-phenylmethanesulfonyl-propionamide;
- (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-(bis-thiophen-2-ylmethyl-amino)-3-phenylmethanesulfonyl-propionamide;
- 10 (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-dibenzylamino-3-phenylmethanesulfonyl-propionamide;
- (S)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-(tetrahydro-pyran-4-ylamino)-3-thiophen-2-yl-propionamide;
- 15 S)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-isopropylamino-3-thiophen-2-yl-propionamide;
- (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-isopropylamino-3-phenylmethanesulfonyl-propionamide;
- (R)-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- 20 R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-[(2-methoxy-ethyl)-(tetrahydro-pyran-4-yl)-amino]-3-phenylmethanesulfonyl-propionamide;
- 25 (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-cyclohexylamino-3-phenylmethanesulfonyl-propionamide;
- (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-dimethylamino-3-phenylmethanesulfonyl-propionamide;
- 30 N-cyanomethyl-3-cyclohexyl-propionamide;
- N-cyanomethyl-3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionamide;
- 3-(3-cyclohexyl-propionylamino)-2-oxo-5-phenyl-pentanoic acid thiazol-2-ylamide;
- 3-cyclohexyl-N-(1-formyl-3-phenyl-propyl)-propionamide;
- 3-(2-difluoromethoxy-phenylmethanesulfonyl)-N-[(S)-1-(5-ethyl-[1,3,4]oxadiazole-2-carbonyl)-propyl]-propionamide;
- 35 N-[(S)-1-(benzoxazole-2-carbonyl)-propyl]-2-(2-cyano-phenylamino)-3-cyclohexyl-propionamide;
- N-Cyanomethyl-3-cyclohexyl-2-(4-methoxy-phenoxy)-propionamide;
- 2-benzyloxy-N-cyanomethyl-3-cyclohexyl-propionamide;

-50-

- (*R*)-*N*-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-butyl]-2-benzyloxy-3-phenylmethanesulfonyl-propionamide;
- (*R*)-*N*-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propyl]-2-methoxymethoxy-3-phenylmethanesulfonyl-propionamide;
- 5 (*S*)-*N*-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-butyl]-2-hydroxy-3-phenyl-propionamide;
- (*R*)-*N*-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propyl]-3-phenylmethanesulfonyl-2-triisopropylsilanyloxy-propionamide;
- (*R*)-*N*-[(*S*)-1-(1-benzothiazol-2-yl-methanoyl)-propyl]-2-hydroxy-3-phenylmethanesulfonyl-propionamide;
- 10 (*R*)-2-hydroxy-3-phenylmethanesulfonyl-*N*-[(*S*)-1-(1-pyridazin-3-yl-methanoyl)-butyl]-propionamide;
- (*S*)-3-((*R*)-2-hydroxy-3-phenylmethanesulfonyl-propanoylamino)-2-oxo-pentanoic acid benzylamide;
- (*R*)-*N*-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propyl]-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionamide;
- 15 (*R*)-*N*-[(*S*)-1-(1-benzothiazol-2-yl-methanoyl)-propyl]-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionamide;
- (2*R*,5*S*)-2-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonylmethyl]-6-ethoxy-5-ethyl-morpholin-3-one; and their corresponding N-oxides, and their prodrugs, and their protected
- 20 derivatives, individual isomers and mixtures of isomers thereof; and the pharmaceutically acceptable salts and solvates (e.g. hydrates) of such compounds and their N-oxides and their prodrugs, and their protected derivatives, individual isomers and mixtures of isomers thereof.

Pharmacology and Utility:

- 25 The compounds of the invention are selective inhibitors of cathepsin S and, as such, are useful for treating diseases in which cathepsin S activity contributes to the pathology and/or symptomatology of the disease. For example, the compounds of the invention may be useful in treating autoimmune disorders, including, but not limited to, juvenile onset diabetes, multiple sclerosis, pemphigus vulgaris, Graves' disease, myasthenia gravis, systemic lupus
- 30 erythematosus, rheumatoid arthritis and Hashimoto's thyroiditis, allergic disorders, including, but not limited to, asthma, and allogeneic immune responses, including, but not limited to, organ transplants or tissue grafts.

Cathepsin S also is implicated in disorders involving excessive elastolysis, such as

chronic obstructive pulmonary disease (e.g., emphysema), bronchiolitis, excessive airway elastolysis in asthma and bronchitis, pneumonitis and cardiovascular disease such as plaque rupture and atheroma. Cathepsin S is implicated in fibril formation and, therefore, inhibitors of cathepsins S are of use in treatment of systemic amyloidosis.

5 The cysteine protease inhibitory activities of the compounds of the invention can be determined by methods known to those of ordinary skill in the art. Suitable *in vitro* assays for measuring protease activity and the inhibition thereof by test compounds are known. Typically, the assay measures protease induced hydrolysis of a peptide based substrate. Details of assays for measuring protease inhibitory activity are set forth in ENZYME ASSAY EXAMPLES, *infra*.

10

Administration and Pharmaceutical Compositions:

In general, compounds of Formula I will be administered in therapeutically effective amounts via any of the usual and acceptable modes known in the art, either singly or in combination with one or more therapeutic agents. A therapeutically effective amount may vary widely depending on the severity of the disease, the age and relative health of the subject, the potency of the compound used and other factors. For example, therapeutically effective amounts of a compound of Formula I may range from about 1 microgram per kilogram body weight ($\mu\text{g/kg}$) per day to about 60 milligram per kilogram body weight (mg/kg) per day, typically from about 1 $\mu\text{g/kg/day}$ to about 20 mg/kg/day . Therefore, a therapeutically effective amount for a 80 kg human patient may range from about 80 $\mu\text{g/day}$ to about 4.8 g /day, typically from about 80 $\mu\text{g/day}$ to about 1.6 g/day. In general, one of ordinary skill in the art, acting in reliance upon personal knowledge and the disclosure of this Application, will be able to ascertain a therapeutically effective amount of a compound of Formula I for treating a given disease.

The compounds of Formula I can be administered as pharmaceutical compositions by one of the following routes: oral, systemic (e.g., transdermal, intranasal or by suppository) or parenteral (e.g., intramuscular, intravenous or subcutaneous). Compositions can take the form of tablets, pills, capsules, semisolids, powders, sustained release formulations, solutions, suspensions, elixirs, aerosols, or any other appropriate composition and are comprised of, in general, a compound of Formula I in combination with at least one pharmaceutically acceptable excipient. Acceptable excipients are non-toxic, aid administration, and do not adversely affect the therapeutic benefit of the active ingredient. Such excipient may be any solid, liquid, semisolid or, in the case of an aerosol composition, gaseous excipient that is generally available to one of skill in the art.

Solid pharmaceutical excipients include starch, cellulose, talc, glucose, lactose, sucrose, gelatin, malt, rice, flour, chalk, silica gel, magnesium stearate, sodium stearate, glycerol monostearate, sodium chloride, dried skim milk, and the like. Liquid and semisolid excipients may be selected from water, ethanol, glycerol, propylene glycol and various oils,

-52-

including those of petroleum, animal, vegetable or synthetic origin (e.g., peanut oil, soybean oil, mineral oil, sesame oil, and the like). Preferred liquid carriers, particularly for injectable solutions, include water, saline, aqueous dextrose and glycols.

The amount of a compound of Formula I in the composition may vary widely depending upon the type of formulation, size of a unit dosage, kind of excipients and other factors known to those of skill in the art of pharmaceutical sciences. In general, a composition of a compound of Formula I for treating a given disease will comprise from 0.01%w to 10%w, preferably 0.3%w to 1%w, of active ingredient with the remainder being the excipient or excipients. Preferably the pharmaceutical composition is administered in a single unit dosage form for continuous treatment or in a single unit dosage form ad libitum when relief of symptoms is specifically required. Representative pharmaceutical formulations containing a compound of Formula I are described in Example 15, *infra*.

Chemistry:

Processes for Making Compounds of Formula I:

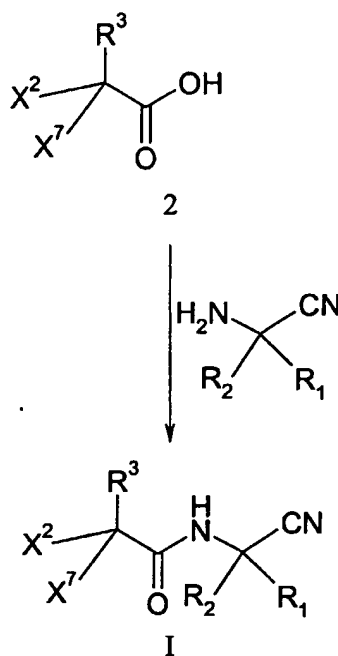
Compounds of the invention may be prepared by the application or adaptation of known methods, by which is meant methods used heretofore or described in the literature, for example those described by R.C. Larock in Comprehensive Organic Transformations, VCH publishers, 1989.

In the reactions described hereinafter it may be necessary to protect reactive functional groups, for example hydroxy, amino, imino, thio or carboxy groups, where these are desired in the final product, to avoid their unwanted participation in the reactions. Conventional protecting groups may be used in accordance with standard practice, for examples see T.W. Greene and P. G. M. Wuts in "Protective Groups in Organic Chemistry" John Wiley and Sons, 1991.

Compounds of Formula I, where X^1 is $-NHC(R^1)(R^2)X^3$, can be prepared by proceeding as in the following Reaction Scheme 1:

Reaction Scheme 1

-53-



in which each X², X³, X⁷, R¹, R² and R³ are as defined for Formula I in the Summary of the Invention.

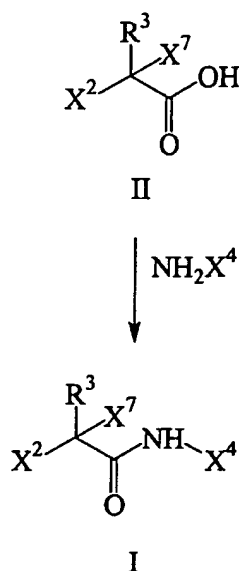
Compounds of Formula I can be prepared by condensing an acid of Formula II with an amino
 5 compound of formula NH₂CR¹R²X³. The condensation reaction can be effected with an appropriate coupling
 agent (e.g., benzotriazol-1-yloxytrispyrrolidinophosphonium hexafluorophosphate (PyBOP[®]), tetra-
 methyluroniumhexafluorophosphate (HATU), 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride
 (EDCI), O-benzotriazol-1-yl-N,N,N',N'-tetramethyluronium hexafluorophosphate (HBTU),
 1,3-dicyclohexylcarbodiimide (DCC), N-cyclohexylcarbodiimide, N'-methylpolystyrene, or the like) and
 10 optionally an appropriate catalyst (e.g., 1-hydroxybenzotriazole (HOBt), 1-hydroxy-7-azabenzotriazole (HOAt),
 O-(7-azabenzotriazol-1-yl)-1,1,3,3, , or the like) and non-nucleophilic base (e.g., triethylamine,
 N-methylmorpholine, and the like, or any suitable combination thereof) at ambient temperature and requires 5 to
 10 hours to complete.

An oxidation step, if required, can be carried out with an oxidizing agent (e.g., Oxone[®],
 15 metachloroperbenzoic acid or the like) in a suitable solvent (e.g., methanol, water, or the like, or any suitable
 combination thereof) at ambient temperature and requires 16 to 24 hours to complete. Detailed descriptions for
 the synthesis of a compound of Formula I by the processes in Reaction Scheme 1 are set forth in the Examples 1
 to 10, *infra*.

Compounds of Formula I, where X¹ is -NHX⁴, can be prepared by proceeding as in the
 20 following Reaction Scheme 2:

Reaction Scheme 2

-54-



in which each X^2 , X^4 , X^7 and R^3 are as defined for Formula I in the Summary of the Invention.

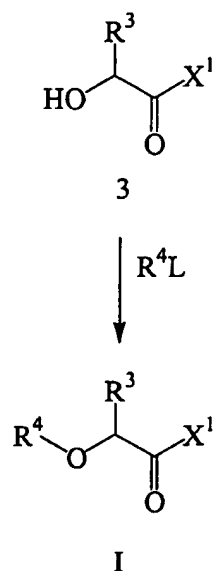
Compounds of Formula I can be prepared by condensing an acid of Formula II with an amino
 5 compound of formula NH_2X^4 . The condensation reaction can be effected with an appropriate coupling agent
 (e.g., benzotriazol-1-yloxytrispyrrolidinophosphonium hexafluorophosphate (PyBOP[®]), *O*-(7-azabenzotriazol-1-
 yl)-1,1,3,3, tetra-methyluroniumhexafluorophosphate (HATU), 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide
 hydrochloride (EDCI), *O*-benzotriazol-1-yl-*N,N,N',N'*-tetramethyluronium hexafluorophosphate (HBTU),
 1,3-dicyclohexylcarbodiimide (DCC), *N*-cyclohexylcarbodiimide, *N'*-methylpolystyrene, or the like) and
 10 optionally an appropriate catalyst (e.g., 1-hydroxybenzotriazole (HOBT), 1-hydroxy-7-azabenzotriazole (HOAt),
 or the like) and non-nucleophilic base (e.g., triethylamine, *N*-methylmorpholine, and the like, or any suitable
 combination thereof) at ambient temperature and requires 5 to 10 hours to complete.

An oxidation step, if required, can be carried out with an oxidizing agent (e.g., Oxone[®],
 metachloroperbenzoic acid or the like) in a suitable solvent (e.g., methanol, water, or the like, or any suitable
 15 combination thereof) at ambient temperature and requires 16 to 24 hours to complete.

Compounds of Formula I in which X^2 is $-\text{OR}^4$, can be prepared by reacting a
 compound of Formula 3 with a compound of Formula R^4L according to the following reaction
 scheme:

Reaction Scheme 3

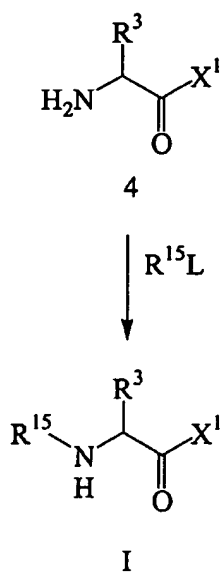
-55-



in which L is a leaving group and X^1 , R^3 and R^4 are as defined in the Summary of the Invention. A detailed description for the synthesis of a compound of Formula I by the process described above is set forth in Example 4, *infra*.

- 5 Compounds of Formula I, in which X^2 is $-\text{NHR}^{15}$, can be prepared by reacting a compound of Formula 4 with a compound of Formula R^{15}L according to the following reaction scheme:

Reaction Scheme 4



10

in which L is a leaving group and X^1 , R^3 and R^{15} are as defined in the Summary of the Invention. A detailed description for the synthesis of a compound of Formula I by the process described above is set forth in [update], *infra*.

Additional Processes for Preparing Compounds of Formula I:

A compound of Formula I can be prepared as a pharmaceutically acceptable acid addition salt by reacting the free base form of the compound with a pharmaceutically acceptable inorganic or organic acid. Alternatively, a pharmaceutically acceptable base addition salt of a compound of Formula I can be prepared by reacting the free acid form of the compound with a pharmaceutically acceptable inorganic or organic base. Inorganic and organic acids and bases suitable for the preparation of the pharmaceutically acceptable salts of compounds of Formula I are set forth in the definitions section of this Application. Alternatively, the salt forms of the compounds of Formula I can be prepared using salts of the starting materials or intermediates.

The free acid or free base forms of the compounds of Formula I can be prepared from the corresponding base addition salt or acid addition salt form. For example, a compound of Formula I in an acid addition salt form can be converted to the corresponding free base by treating with a suitable base (e.g., ammonium hydroxide solution, sodium hydroxide, and the like). A compound of Formula I in a base addition salt form can be converted to the corresponding free acid by treating with a suitable acid (e.g., hydrochloric acid, etc).

The *N*-oxides of compounds of Formula I can be prepared by methods known to those of ordinary skill in the art. For example, *N*-oxides can be prepared by treating an unoxidized form of the compound of Formula I with an oxidizing agent (e.g., trifluoroperacetic acid, permaleic acid, perbenzoic acid, peracetic acid, *meta*-chloroperoxybenzoic acid, or the like) in a suitable inert organic solvent (e.g., a halogenated hydrocarbon such as dichloromethane) at approximately 0°C. Alternatively, the *N*-oxides of the compounds of Formula I can be prepared from the *N*-oxide of an appropriate starting material.

Compounds of Formula I in unoxidized form can be prepared from *N*-oxides of compounds of Formula I by treating with a reducing agent (e.g., sulfur, sulfur dioxide, triphenyl phosphine, lithium borohydride, sodium borohydride, phosphorus trichloride, tribromide, or the like) in an suitable inert organic solvent (e.g., acetonitrile, ethanol, aqueous dioxane, or the like) at 0 to 80°C.

Prodrug derivatives of the compounds of Formula I can be prepared by methods known to those of ordinary skill in the art (e.g., for further details see Saulnier *et al.* (1994), *Bioorganic and Medicinal Chemistry Letters*, Vol. 4, p. 1985). For example, appropriate prodrugs can be prepared by reacting a non-derivatized compound of Formula I with a

suitable carbamylating agent (e.g., 1,1-acyloxyalkylcarbonochloridate, *para*-nitrophenyl carbonate, or the like).

Protected derivatives of the compounds of Formula I can be made by means known to those of ordinary skill in the art. A detailed description of the techniques applicable to the creation of protecting groups and their removal can be found in T.W. Greene, *Protecting Groups in Organic Synthesis*, 3rd edition, John Wiley & Sons, Inc. 1999.

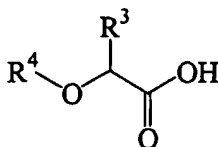
Compounds of the present invention may be conveniently prepared, or formed during the process of the invention, as solvates (e.g. hydrates). Hydrates of compounds of the present invention may be conveniently prepared by recrystallisation from an aqueous/organic solvent mixture, using organic solvents such as dioxin, tetrahydrofuran or methanol.

Compounds of Formula I can be prepared as their individual stereoisomers by reacting a racemic mixture of the compound with an optically active resolving agent to form a pair of diastereoisomeric compounds, separating the diastereomers and recovering the optically pure enantiomer. While resolution of enantiomers can be carried out using covalent diasteromeric derivatives of compounds of Formula I, dissociable complexes are preferred (e.g., crystalline diastereoisomeric salts). Diastereomers have distinct physical properties (e.g., melting points, boiling points, solubilities, reactivity, etc.) and can be readily separated by taking advantage of these dissimilarities. The diastereomers can be separated by chromatography or, preferably, by separation/resolution techniques based upon differences in solubility. The optically pure enantiomer is then recovered, along with the resolving agent, by any practical means that would not result in racemization. A more detailed description of the techniques applicable to the resolution of stereoisomers of compounds from their racemic mixture can be found in Jean Jacques Andre Collet, Samuel H. Wilen, *Enantiomers, Racemates and Resolutions*, John Wiley & Sons, Inc. (1981).

25

In summary, the compounds of Formula I are made by a process which comprises:

(A) reacting a compound of Formula II:



II

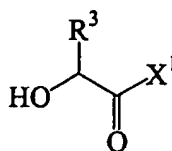
with a compound of the formula $\text{NH}_2\text{CR}^1\text{R}^2\text{X}^3$, in which X^3 , R^1 , R^2 , R^3 and R^4 are as defined in the Summary of the Invention for Formula I; or

30

-58-

(B) reacting a compound of Formula II with a compound of the formula NH_2X^4 , in which X^4 , R^3 and R^4 are as defined in the Summary of the Invention for Formula I; or

(C) reacting a compound of Formula 3:

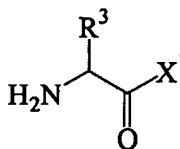


3

5

with a compound of formula R^4L , in which X^1 , R^3 and R^4 are as defined in the Summary of the Invention and L is a leaving group; or

10 (D) reacting a compound of Formula 4:



4

with a compound of formula R^{15}L , in which X^1 , R^3 and R^4 are as defined in the Summary of the Invention and L is a leaving group; and

15 (E) optionally converting a compound of Formula I into a pharmaceutically acceptable salt;

(F) optionally converting a salt form of a compound of Formula I to non-salt form;

(G) optionally converting an unoxidized form of a compound of Formula I into a pharmaceutically acceptable *N*-oxide;

(H) optionally converting an *N*-oxide form of a compound of Formula I to its unoxidized form;

20 (I) optionally resolving an individual isomer of a compound of Formula I from a mixture of isomers;

(J) optionally converting a non-derivatized compound of Formula I into a pharmaceutically prodrug derivative; and

(K) optionally converting a prodrug derivative of a compound of Formula I to its non-derivatized form.

Examples:

25 The present invention is further exemplified, but not limited by, the following examples that illustrate the preparation of compounds of Formula I and II (Examples) and intermediates (References) according to the invention.

LC/MS-Procedures:

30 LC/MS (Method A):

Mass Spectrometer (MS) - LCT Time-of-Flight (Micromass UK Ltd) Serial No. KA014

-59-

Ionization Mode: Electrospray (Positive Ion)

Scan: ToF MS (Full Scan m/z 100 - 1200, sum for 0.4 s @ 50us/scan) Centroid Mode

Liquid Chromatograph (LC): Hewlett Packard HP1100 Series Binary Pump (Serial # US80301343)
& Degasser (serial # JP73008973)

5 Mobile Phase:

A = Water + 0.05% TFA (trifluoroacetic acid) buffer

B = Acetonitrile + 0.05% TFA buffer

Gradient: 5%B to 100%B in 5 minutes

Column: Hypersil BDS C-18, 3u, 4.6mm x 50mm Reverse Phase

10 Injection volume: 5 uL

Flow rate: 1ml/min to column & to UV detector, flow split after UV detector
such that 0.75ml/min to ELS detector and 0.25ml/min to mass spectrometer.

Auxiliary Detectors: (i) Hewlett Packard Model HP1100 Series UV detector (serial # JP73704703)
15 wavelength = 220nm

(ii) Sedere (France) Model SEDEX 75 Evaporative Light Scattering (ELS) detector
(serial # 9970002A)

temperature = 46 deg C, Nitrogen pressure = 4bar

20 Autosampler / Injector: Gilson Model 215 Liquid Handler with Model 819 injection valve (serial #
259E8280)

LC/MS (Method B):

Same as method A, but with a different gradient: 5%B to 90%B in 3 minutes, 90%B to 100%B in 2
25 min

LC/MS (Method C):

Mass Spectrometer (MS) - LCT Time-of-Flight (Micromass UK Ltd) Serial No. KA014

Ionization Mode: Electrospray (Positive Ion)

30 Scan: ToF MS (Full Scan m/z 100 - 1200, sum for 0.4 s @ 50us/scan) Centroid Mode

Liquid Chromatograph (LC): Hewlett Packard HP1100 Series Binary Pump (Serial # US80301343)
& Degasser (serial # JP73008973)

Mobile Phase:

A = Water + 0.1% formic acid buffer

35 B = Acetonitrile + 0.1% formic acid buffer

Gradient: 5%B to 90%B in 3 minutes, 90%B to 100%B in 2 min

-60-

Column: Phenomenex Synergi C-18, 2u, 4mm x 20mm Reverse Phase

Injection volume: 5 uL

Flow rate: 1ml/min to column & to UV detector, flow split after UV detector
such that 0.75ml/min to ELS detector and 0.25ml/min to mass spectrometer.

5

Auxiliary Detectors: (i) Hewlett Packard Model HP1100 Series UV detector (serial # JP73704703)
wavelength = 220nm

(ii) Sedere (France) Model SEDEX 75 Evaporative Light Scattering (ELS) detector
(serial # 9970002A)

10

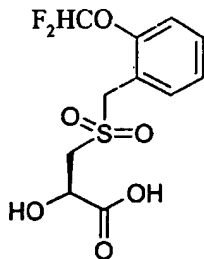
temperature = 46 deg C, Nitrogen pressure = 4bar

Autosampler / Injector: Gilson Model 215 Liquid Handler with Model 819 injection valve (serial #
259E8280)

15

REFERENCE EXAMPLE 1

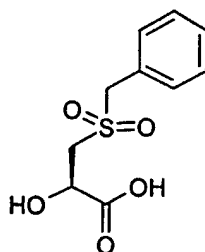
(a) (R)-3-[2-(1,1-Difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionic acid



A solution of (R)-2-*tert*-Butoxycarbonylamino-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-
propionic acid (5.19g) in CH₂Cl₂ (20mL), was treated with trifluoroacetic acid (20mL) at room
20 temperature. After two hours, the reaction mixture was concentrated under reduced pressure. The
white solid obtained was dissolved in 1M H₂SO₄ (100mL) and dioxane (30mL). The solution was
cooled to 0°C, NaNO₂ (1.95g in 50mL of water) was added with stirring for 1 hour. The reaction
mixture was stirred overnight at ambient temperature. The product was then concentrated and
extracted with ethyl acetate, dried with anhydrous MgSO₄, filtered, concentrated and recrystallized
25 from ethyl acetate to yield (R)-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-
propionic acid (2.36g).

(b) (R)-2-hydroxy-3-phenylmethanesulfonyl-propionic acid

-61-



By proceeding in a manner similar to Reference Example 1(a) above but using (R)-2-*tert*-butoxycarbonylamino-3-[phenylmethanesulfonyl]-propionic acid there was prepared (R)-2-hydroxy-3-phenylmethanesulfonyl-propionic acid.

5

REFERENCE EXAMPLE 2

(R)-2-Amino-N-methoxy-N-methyl-butylamide

To a solution of [(R)-1-(methoxy-methyl-carbamoyl)-propyl]-carbamic acid *tert*-butyl ester (4.92g, 20mmol) in CH₂Cl₂ (20ml) was added TFA (10mL) at room temperature. After stirring for 2 hours, the reaction mixture was concentrated to dryness under reduced pressure to produce (R)-2-amino-N-methoxy-N-methyl-butylamide TFA salt (5.4g).

10

REFERENCE EXAMPLE 3

(R)-3-[2-(1,1-Difluoro-methoxy)-phenylmethanesulfonyl]-2-triisopropylsilanyloxy-propionic acid

(R)-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionic acid (7.0g, 22.58mmol), in CH₂Cl₂ (50mL) was reacted with 2, 6-lutidine (12.09g, 112.9mmol) and triisopropylsilyl-trifluoro-methanesulfonate (20.75g, 67.74mmol) at -78°C for one hour. The reaction mixture was allowed to warm to room temperature before being quenched by the addition of saturated ammonium chloride solution. The product was extracted with ethyl acetate, the solvent was removed under reduced pressure and the oil residue was then dissolved in EtOH:THF:H₂O (3:1:1, 60mL). Solid K₂CO₃ (24g) was added at room temperature and the mixture was stirred for one hour, filtered, extracted with ethyl acetate, dried with anhydrous MgSO₄, filtered and concentrated to yield (R)-3-[2-(1,1-Difluoro-methoxy)-phenylmethanesulfonyl]-2-triisopropylsilanyloxy-propionic acid (8.58g).

20

25

Following as in reference 3 provided the following intermediate:

(R)-3-Phenylmethanesulfonyl-2-triisopropylsilanyloxy-propionic acid

30

REFERENCE EXAMPLE 4

3-[2-(1,1-Difluoro-methoxy)-phenylmethanesulfonyl]-propionic acid

-62-

A mixture of [2-(1,1-difluoro-methoxy)-phenyl]-methanethiol (190mg, 1.0mmol), acrylic acid (69μL, 1.0mmol), diisopropylethylamine (440 μL, 1.1mmol) and 0.5mL dimethylformamide was stirred at 45°C for 4 hours. Diethyl ether (5mL) and 1N HCl (2mL) was added. The layers were separated and the organic layer was washed with 1N HCl (2mL), dried over MgSO₄ and concentrated. The resulting oil was dissolved in methanol (5mL), treated with an aqueous solution (5mL) of Oxone® (921mg, 1.5mmol), and stirred for 1 hour. Methanol was removed under reduced pressure and 20mL water was added. The mixture was extracted with two 60mL portions of ethyl acetate, dried over MgSO₄ and concentrated to give 3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-propionic acid (160mg; 0.54mmol, 54% yield).

REFERENCE EXAMPLE 5

3-Benzylsulfanyl-2-(2-nitro-phenylamino)-propionic acid

S-benzylcysteine (1.06g, 5.0mmol), 2-fluoronitrobenzene (1.05mL, 10.0mmol), potassium carbonate (1.38g, 10.0mmol) and dimethylformamide (3mL) were combined and stirred at 100°C for 4 hours. The mixture was diluted with 40mL water and washed with two 15mL portions of diethyl ether. The aqueous layer was acidified to pH 4 with 6N HCl and extracted with two 30mL portions of ethyl acetate. The ethyl acetate layer was dried over MgSO₄, and concentrated. Diethyl ether was added and then decanted to give 3-benzylsulfanyl-2-(2-nitro-phenylamino)-propionic acid (541mg, 1.63mmol, 33%yield).

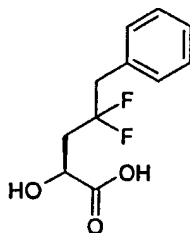
REFERENCE EXAMPLE 6

(R)-3-Benzylsulfanyl-2-(5-nitro-thiazol-2-ylamino)-propionic acid

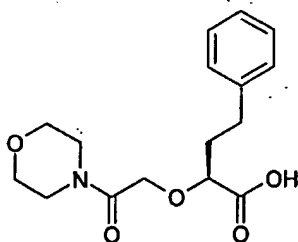
S-benzylcysteine (0.845g, 4mmol) and bis(trimethylsilyl)acetamide (3mL, 16mmol) were stirred at 75°C for 1 hour. 2-Bromo-5-nitrothiazole (837mg, 4mmol) and toluene (8mL) was added and the mixture was stirred at 100°C for 1 day. Toluene was removed under reduced pressure. The residue was stirred in 5mL dioxane and 5mL 1N HCl for 30 minutes. Dioxane was removed under reduced pressure and the mixture was basified with saturated NaHCO₃ and washed with 50mL ethyl acetate. The aqueous layer was acidified with 6N HCl and extracted with two 25mL portions of ethyl acetate, dried over MgSO₄, concentrated and chromatographed using a gradient of 5-10% methanol in methylene chloride to yield (R)-3-benzylsulfanyl-2-(5-nitro-thiazol-2-ylamino)-propionic acid (42.7mg, 0.123mmol, 3% yield).

REFERENCE EXAMPLE 7

-63-

(2S)-4,4-Difluoro-2-hydroxy-5-phenyl-pentanoic acid

To a suspension of (S)-2-Amino-4,4-difluoro-5-phenyl-pentanoic acid (1.0 mmol, 230mg) in water (3mL) was added 2M sulfuric acid dropwise until the solid dissolved (ca 3mL). A solution of sodium nitrite (1.5 eq., 1.5 mmol, 104mg) in 1 ml of water was then added dropwise. The mixture was stirred at room temperature for 21 hours then extracted twice with ether (30 ml). The organic layers were dried over MgSO₄ and then concentrated in vacuum to afford (2S)-4,4-difluoro-2-hydroxy-5-phenyl-pentanoic acid (90 mg, 39%) as a white solid. ¹H NMR (CDCl₃) 7.3 (m, 5H), 5.6 (b, 1H), 4.61 (dd, J=8.5, 2.9 Hz, 1H), 3.3 (t, J=16.8 Hz, 2H), 2.45 (m, 1H), 2.2 (m, 2H).

REFERENCE EXAMPLE 82-(S)-(2-Morpholin-4-yl-2-oxo-ethoxy)-4-phenyl-butyric acid

Step (i): To a cooled (0°C) solution of ethyl (2R) 2-hydroxy-4-phenylbutyrate (1.81g, 8.71 mmol), 4-nitro-benzoic acid (1.1eq., 9.56 mmol, 1.598g) and triphenyl phosphine (1.1 eq., 9.5 mmol, 2.50g) in dry THF (80mL) under nitrogen was added slowly diethyl azodicarboxylate (1.1 eq., 9.56 mmol, 1.67g). The mixture was stirred at 0°C for 2.5 hours and then concentrated in vacuum. The residue was triturated with a mixture of ethyl acetate and heptane (1:3, 150mL) and the resulting solids were filtered off. The filtrate was concentrated in vacuum and purified over 110g silica gel, eluting with a mixture of ethyl acetate and heptane (1:4, v/v) to afford 4-nitro-benzoic acid (S)-1-ethoxycarbonyl-3-phenyl-propyl ester (3.4g, 98%).

Step (ii): To a cooled (0°C) solution of 4-nitro-benzoic acid (S)-1-ethoxycarbonyl-3-phenyl-propyl ester (2.04 g, 5.83 mmol) in MeOH (30 mL) was added potassium carbonate (1.5 eq., 8.75 mmol, 1.21g). The mixture was stirred at 0°C for 5 minutes then at room temperature for 1.5 hours and concentrated in vacuum. The residue was partitioned between water (40mL) and ethyl acetate (40mL). The organic layer was dried over MgSO₄ and then concentrated in vacuum. The residue was purified

-64-

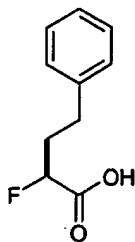
over 35g silica gel, eluted with dichloromethane to afford methyl-(2S)-2-hydroxy-4-phenyl-butyrate as a colorless oil (933mg, 82%).

Step (iii): To a solution of methyl-(2S)-2-hydroxy-4-phenyl-butyrate (300mg, 1.54 mmol) in dry DMF (3mL) under nitrogen was added sodium hydride (60%, 1.5 eq., 2.32 mmol, 92.7mg). After 5 min, 4-(2-chloroacetyl) morpholine (1.1 eq., 1.69 mmol, 277mg) was added and the mixture was stirred at room temperature for 24 hours, then diluted with water (60mL) and then neutralized with 1 N HCl. The aqueous solution was extracted twice with ethyl acetate (40mL). The organic layer was washed with water (50mL), dried over MgSO₄ and then concentrated in vacuum. The residue was purified over 35g silica gel, eluting with ethyl acetate then with 5% MeOH in ethyl acetate to afford (S)-2-(2-morpholin-4-yl-2-oxo-ethoxy)-4-phenyl-butyric acid methyl ester (117mg, 24%).

Step (iv): To a solution of (S)-2-(2-morpholin-4-yl-2-oxo-ethoxy)-4-phenyl-butyric acid methyl ester (117mg, 0.36 mmol) in MeOH:H₂O (2:1 vol, 3mL) was added lithium hydroxide hydrate (2.0 eq., 0.73 mmol, 30.5mg). The mixture was stirred at room temperature for 5 hours, then diluted with water (30mL) and then extracted with ether (30mL). The aqueous layer was acidified with 1N HCl and extracted twice with ether (30mL). The acidic extracts were dried over MgSO₄ and then concentrated in vacuum to afford (S)-2-(2-Morpholin-4-yl-2-oxo-ethoxy)-4-phenyl-butyric acid (85.5mg, 77%) as a colorless oil. ¹H NMR (CDCl₃) 10.5 (b, 1H), 7.2 (m, 5H), 4.55 (d, J=15.2 Hz, 1H), 4.14 (d, J=15.2 Hz, 1H), 3.9 (dd, J=7.6, 4.2 Hz, 1H), 4.6 (m, 6H), 3.4 (m, 2H), 2.8 (m, 2H), 2.3 (m, 1H), 2.15 (m, 1H). LC/MS 96% (M+1) 308.

REFERENCE EXAMPLE 9

(2S)-2-Fluoro-4-phenyl-butyric acid



25

Step (i): To a cooled (0°C) solution of methyl-(2R)-2-hydroxy-4-phenyl-butyrate (1.00g, 4.80 mmol) in dry dichloromethane (3mL) was added DAST (3.0eq., 14.4 mmol, 2.32g). The mixture was stirred at room temperature for 18 hours then diluted with dichloromethane (20mL) and carefully quenched with saturated sodium bicarbonate (150mL). The aqueous layer was extracted with dichloromethane (30mL) and the organic layers were dried over MgSO₄ and then concentrated in vacuo. The residue

30

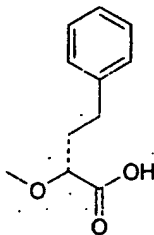
-65-

was purified over 90g silica gel, eluting with a mixture of dichloromethane and heptane (1:2 then 1:1, v/v) to afford methyl-2S-fluoro-4-phenyl-butyrate as a light yellow oil (578 mg, 57%).

Step (ii): To a solution of methyl-2S-fluoro-4-phenyl-butyrate (577mg, 2.74 mmol) in a mixture of MeOH:H₂O (2:1 vol, 6mL) was added lithium hydroxide monohydrate (1.5 eq., 4.11 mmol, 173mg). The mixture was stirred at room temperature for 5 hours and then concentrated in vacuum. The residue was diluted with water (30mL) and extracted with ether (20mL). The aqueous layer was acidified with HCl and extracted with ether (30mL). The acidic extract was dried over MgSO₄ and then concentrated in vacuum to afford 2(S)-fluoro-4-phenyl-butyric acid as a yellow oil (486 mg, 97%). ¹H NMR (CDCl₃) 7.5 (b, 1H), 7.3 (m, 5H), 4.95 (ddd, J=48.9, 6.9, 5.4 Hz, 1H), 2.85 (m, 2H), 2.25 (m, 2H). MS (CI) M+1 183.

REFERENCE EXAMPLE 10

2(R)-Methoxy-4-phenyl-butyric acid



15

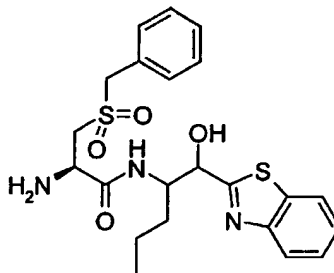
Step 1: To a solution of ethyl-(2R)-2-hydroxy-4-phenyl-butyrate (500mg, 2.40 mmol) in dry DMF (4mL) under nitrogen was added sodium hydride (60%, 2.0 eq., 4.80 mmol, 192mg) followed by methyl iodide (3.0 eq., 7.20 mmol, 1.02g). The mixture was stirred at room temperature for 22 hours, then diluted with NH₄Cl (100mL) and extracted with ethyl acetate (50mL). The organic layer was dried over MgSO₄ and then concentrated in vacuum. The residue was purified over 35g silica gel, eluting with ethyl acetate and heptane (1:3, v/v) to afford (R)-2-methoxy-4-phenyl-butyric acid ethyl ester (480 mg, 90%).

Step 2: To a solution of (R)-2-methoxy-4-phenyl-butyric acid ethyl ester (480mg, 2.8 mmol) in MeOH:H₂O (2:1 vol, 9mL) was added lithium hydroxide hydrate (2.0 eq., 4.32 mmol, 181mg). The mixture was stirred at room temperature for 2.5 hours, then diluted with water (20mL) and then extracted with ether (20mL). The aqueous layer was acidified with 1N HCl and then extracted twice with ether (30 mL). The combined extracts were dried over MgSO₄ and then concentrated in vacuum to afford 2(R)-methoxy-4-phenyl-butyric acid (426mg, quant.) as a colorless solid. ¹H NMR (CDCl₃) 7.25 (m, 5H), 3.8 (dd, J=6.8, 5.2 Hz, 1H), 3.48 (s, 3H), 2.78 (t, J=7.3 Hz, 2H), 2.1 (m, 2H). MS (CI) M 194.

Following as in reference 10 but using benzyl bromide in step 2 provided the following intermediate:
2(R)-Benzyloxy-4-phenyl-butyric acid

REFERENCE EXAMPLE 11

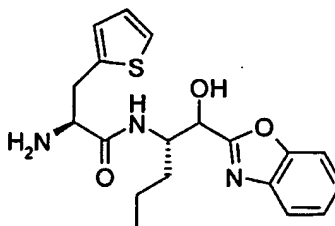
- 5 (a) (R)-2-Amino-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-propionamide



A solution of {(R)-1-[1-(benzothiazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester {888mg, 1.58mmol, Example 27(a)} in dichloromethane (5mL) was treated with trifluoroacetic acid (5mL). The mixture was stirred at room temperature for one hour and then evaporated. The residue was dissolved in dichloromethane (20mL) and this solution was treated with Silicycle Triamine (4.3g, 16mmol). The mixture was stirred at room temperature for two hours and then filtered. The filtrate was evaporated to give the title compound (692mg, 94%). LC/MS m/z=562 (M+H).

15

- (b) (S)-2-Amino-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-thiophen-2-yl-propionamide

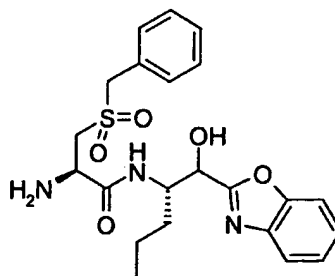


By proceeding in a manner similar to Reference Example 11(a) above but using {(S)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-thiophen-2-yl-ethyl}-carbamic acid tert-butyl ester {790mg, 1.67mmol, Example 27(c)} and subjecting the reaction product to flash chromatography on silica eluting with a mixture of ethyl acetate and methanol (9:1, v/v) there was prepared (S)-2-amino-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-thiophen-2-yl-propionamide (415mg, 66%). LC/MS m/z=374 (M+H).

25

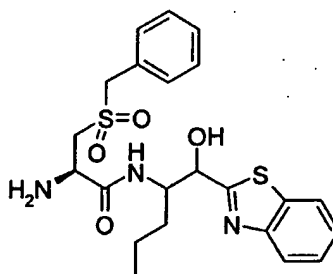
- (c) (R)-2-Amino-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-

-67-

propionamide

By proceeding in a manner similar to Reference Example 11(a) above but using {(R)-1-[(S)-1-
 5 (Benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid
 tert-butyl ester {908mg, 1.66mmol, Example 27(b)} there was prepared (R)-2-amino-N-[(S)-1-
(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-propionamide (726mg, 98%).
 LC/MS $m/z=446$ (M+H).

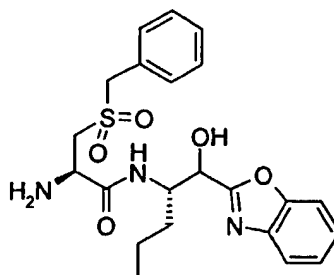
- 10 (d) (R)-2-Amino-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-
propionamide



By proceeding in a manner similar to Reference Example 11(a) above but using {(R)-1-[1-
 (Benzothiazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid
 15 tert-butyl ester {0.63mmol, Example 27(d)} there was prepared (R)-2-Amino-N-[1-(benzothiazol-2-yl-
hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-propionamide (212mg, 73%). LC/MS $m/z=462$
 (M+H).

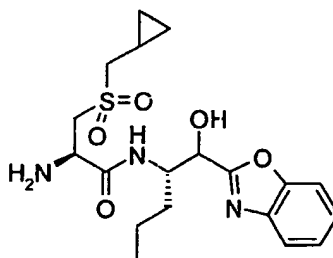
- (e) (R)-2-Amino-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-
 20 propionamide

-68-



By proceeding in a manner similar to Reference Example 11(a) above but using {(R)-1-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester {1.7mmol, Example 27(e)} there was prepared (R)-2-amino-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-propionamide (726mg, 98%). LC/MS $m/z=446$ (M+H).

(f) (R)-2-Amino-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-cyclopropylmethanesulfonyl-propionamide

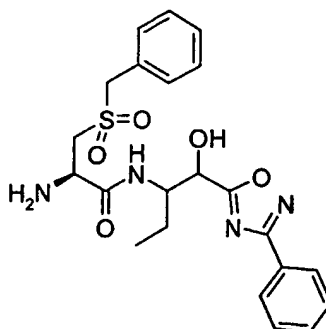


By proceeding in a manner similar to Reference Example 11(a) above but using {(R)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-cyclopropylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester {450mg, 0.88mmol, Example 27(f)} there was prepared (R)-2-amino-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-cyclopropylmethanesulfonyl-propionamide (360mg, 0.879mmol, 100%).

LC/MS $m/z=410$ (M+H)

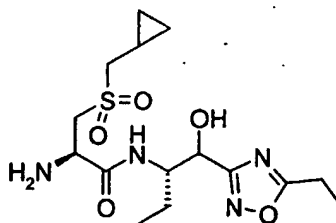
(g) (R)-2-Amino-N-{1-[hydroxy-(3-phenyl-1,2,4-oxadiazol-5-yl)-methyl]-propyl}-3-phenylmethanesulfonyl-propionamide

-69-



By proceeding in a manner similar to Reference Example 11(a) above but using (R)-1-{1-[Hydroxy-(3-phenyl-1,2,4-oxadiazol-5-yl)-methyl]-propylcarbamoyl}-2-phenylmethanesulfonyl-ethyl)-carbamic acid tert-butyl ester {Example 27(g)} there was prepared (R)-2-amino-N-{1-[hydroxy-(3-phenyl-1,2,4-oxadiazol-5-yl)-methyl]-propyl}-3-phenylmethanesulfonyl-propionamide. LC/MS $m/z=481$ (M+Na), 459(M+H)

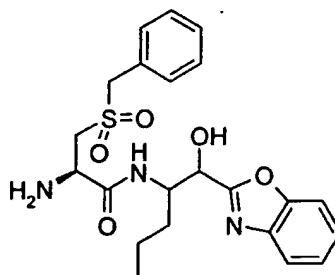
(h) (R)-2-Amino-3-cyclopropylmethanesulfonyl-N-[(S)-1-[(5-ethyl-1,2,4-oxadiazol-3-yl)-hydroxy-methyl]-propyl]-propionamide



By proceeding in a manner similar to Reference Example 11(a) above but using ((R)-2-cyclopropylmethanesulfonyl-1-[(S)-1-[(5-ethyl-1,2,4-oxadiazol-3-yl)-hydroxy-methyl]-propylcarbamoyl]-ethyl)-carbamic acid tert-butyl ester {Example 27(i)} there was prepared (R)-2-amino-3-cyclopropylmethanesulfonyl-N-[(S)-1-[(5-ethyl-1,2,4-oxadiazol-3-yl)-hydroxy-methyl]-propyl]-propionamide. LC/MS $m/z=375$ (M+H)

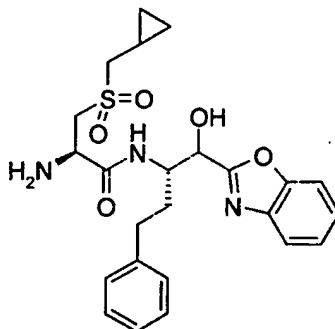
(i) (R)-2-Amino-N-[1-(benzoxazol-2-yl)-hydroxy-methyl]-butyl]-3-phenylmethanesulfonyl-propionamide

-70-



By proceeding in a manner similar to Reference Example 11(a) above but using {(R)-1-[1-(Benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester {Example 27(j)} there was prepared (R)-2-Amino-N-[1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-propionamide. LC/MS $m/z=446(M+H)$

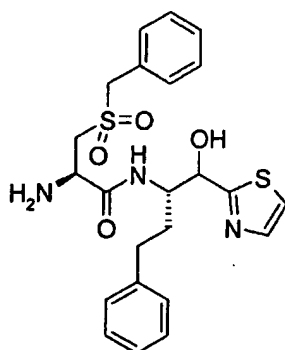
(j) (R)-2-Amino-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-3-phenyl-propyl]-3-cyclopropylmethanesulfonyl-propionamide



By proceeding in a manner similar to Reference Example 11(a) above but using {(R)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-3-phenyl-propylcarbamoyl]-2-cyclopropylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester {Example 27(k)} there was prepared (R)-2-amino-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-3-phenyl-propyl]-3-cyclopropylmethanesulfonyl-propionamide. LC/MS $m/z=472(M+H)$

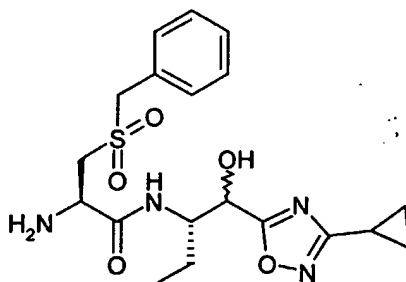
(k) (R)-2-Amino-N-[(S)-1-(hydroxy-thiazol-2-yl-methyl)-3-phenyl-propyl]-3-phenylmethanesulfonyl-propionamide

-71-



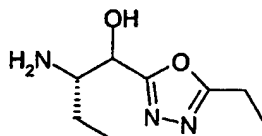
By proceeding in a manner similar to Reference Example 11(a) above but using {(R)-1-[(S)-1-(Hydroxy-thiazol-2-yl-methyl)-3-phenyl-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester {Example 27(l)} there was prepared (R)-2-amino-N-[(S)-1-(hydroxy-thiazol-2-yl-methyl)-3-phenyl-propyl]-3-phenylmethanesulfonyl-propionamide.

(l) (R)-2-Amino-3-phenylmethanesulfonyl-N-[(S)-1-[(3-cyclopropyl-1,2,4-oxadiazol-5-yl)-hydroxy-methyl]-propyl]-propionamide



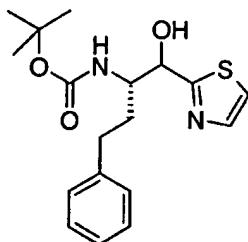
By proceeding in a manner similar to Reference Example 11(a) above but using ((R)-2-phenylmethanesulfonyl-1-[(S)-1-[(3-cyclopropyl-1,2,4-oxadiazol-5-yl)-hydroxy-methyl]-propylcarbamoyl]-ethyl)-carbamic acid tert-butyl ester {Example 27(s)} there was prepared (R)-2-amino-3-phenylmethanesulfonyl-N-[(S)-1-[(3-cyclopropyl-1,2,4-oxadiazol-5-yl)-hydroxy-methyl]-propyl]-propionamide.

(m) 2-amino-1-(5-ethyl-[1,3,4]oxadiazol-2-yl)-butan-1-ol

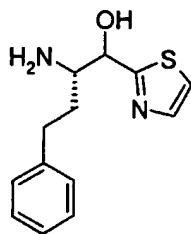


By proceeding in a manner similar to Reference Example 11(a) above but using {1-[(5-ethyl-[1,3,4]oxadiazol-2-yl)-hydroxy-methyl]-propyl}-carbamic acid *tert*-butyl ester (Reference Example 16) there was prepared 2-amino-1-(5-ethyl-[1,3,4]oxadiazol-2-yl)-butan-1-ol.

-72-

REFERENCE EXAMPLE 12[(S)-1-(Hydroxy-thiazol-2-yl-methyl)-3-phenyl-propyl]-carbamic acid tert-butyl ester

n-Butyllithium (4.2ml, 10.5mmol, 2.5M solution in hexanes) was mixed with 16ml diethylether and the resulting solution cooled to -78°C . 2-Bromothiazole (1.64g, 10mmol) was dissolved in a mixture of 2ml diethylether and 1ml THF. This solution was added dropwise to the n-butyllithium solution. The resulting reaction mixture was stirred for 15min. A solution of [(S)-1-(Methoxy-methyl-carbamoyl)-3-phenyl-propyl]-carbamic acid tert-butyl ester (1.4g, 4.3mmol) in 20ml THF was added dropwise to the reaction mixture. Stirring was continued for one hour and the reaction mixture quenched by addition of 50ml water. After warming to room temperature the phases were separated and the aqueous phase extracted with ethyl acetate. The combined organic phases were washed with brine and dried with magnesium sulfate. The solvents were evaporated under vacuum to give 1.4g [(S)-3-Phenyl-1-(thiazole-2-carbonyl)-propyl]-carbamic acid tert-butyl ester as a brown solid. [(S)-3-Phenyl-1-(thiazole-2-carbonyl)-propyl]-carbamic acid tert-butyl ester (1.41g, 4.1mmol) was dissolved in 50 ml ethanol and the solution cooled to 0°C . Sodium borohydride (155mg, 4.1mmol) was added and the reaction mixture stirred for 90 minutes. Water was added and the aqueous phase acidified by addition of 1M hydrochloric acid. The aqueous phase was extracted with ethyl acetate. The combined organic phases were washed with brine and dried with magnesium sulfate. The solvent was evaporated under reduced pressure. (1.32, 3.8mmol, 88%). LC/MS $m/z=271$ (M+H-isobutene), 249(M+H-boc)

REFERENCE EXAMPLE 13(S)-2-Amino-4-phenyl-1-thiazol-2-yl-butan-1-ol

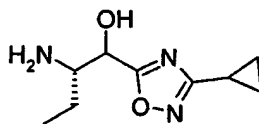
[(S)-1-(Hydroxy-thiazol-2-yl-methyl)-3-phenyl-propyl]-carbamic acid tert-butyl ester (1.32g, 3.8mmol, Reference Example 12) was dissolved in 10ml dichloromethane. Trifluoroacetic acid was added and the resulting reaction mixture stirred for two hours. The solvents were evaporated under

-73-

reduced pressure and saturated sodium bicarbonate solution was added. The solution was extracted with ethyl acetate. The combined organic phases were washed with brine and dried with magnesium sulfate. The solvent was evaporated and the crude product purified via flash chromatography (eluted with ethyl acetate followed by 10% methanol in ethyl acetate) to give (S)-2-amino-4-phenyl-1-thiazol-2-yl-butan-1-ol (466mg, 1.87mmol, 49%). LC/MS $m/z=249(M+H)$.

REFERENCE EXAMPLE 14

(S)-2-Amino-1-(3-cyclopropyl-1,2,4-oxadiazol-5-yl)-butan-1-ol



A solution of boc-3S-amino-2-hydroxypentanoic acid (2.00g, 8.57mmol) and 1.20 equivalents of cyclopropanecarboxamidoxime (1.03g, 10.29mmol) in 20 mL of dichloromethane was stirred at 0°C as 1.25 equivalents of N-cyclohexylcarbodiimide-N'-methyl polystyrene (1.70mmol/g, 6.30g, 10.72mmol) was added in portions and the reaction mixture stirred under nitrogen for three hours while warming to 15°C. The reaction mixture was filtered and the resin washed with dichloromethane. Evaporate under vacuum to dryness. [LC/MS $m/z=338(M+H+Na)$] The residue is dissolved in 20 mL of tetrahydrofuran and heated in a microwave reactor at 160°C for three minutes. Evaporate under vacuum to dryness. [LC/MS $m/z=320(M+H+Na)$] The residue is dissolved in 50 mL of dichloromethane and stirred at room temperature as a 50 mL solution of 50% trifluoroacetic acid in dichloromethane was added dropwise. After three hours the reaction was evaporated under vacuum to dryness and dissolved in 50 mL of dichloromethane again. Three equivalents of Silicycle triamine-3 was added and the mixture stirred at room temperature overnight. The mixture was filtered and washed with dichloromethane. Evaporate under vacuum to give (S)-2-Amino-1-(3-cyclopropyl-1,2,4-oxadiazol-5-yl)-butan-1-ol 1.04g (61% overall). [LC/MS $m/z=198(M+H)$]

REFERENCE EXAMPLE 15

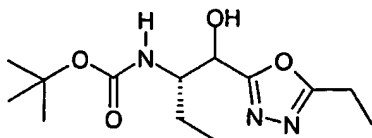
Ethyl-1,3,4-oxadiazole:

A mixture of the formic hydrazide (60g, 1mole), triethylorthopropionate (176.26g, 1mole) and p-toluenesulfonic acid (250mg) was heated at 120°C for 12 hours. The ethanol was removed under vacuum and the residue was distilled under vacuum to yield 24g of ethyl-1,3,4-oxadiazole. H^1 NMR (DMSO- d_6): 9.34 (1H, s), 2.86 (2H, q), 1.25(3H, t).

REFERENCE EXAMPLE 16

{1-[(5-Ethyl-[1,3,4]oxadiazol-2-yl)-hydroxy-methyl]-propyl}-carbamic acid tert-butyl ester

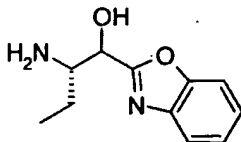
-74-



To a stirred solution of the ethyl-1,3,4-oxadiazole (4.66g, 48mmol, Reference Example 15) in THF (50ml) was added n-BuLi (1.6M solution in 30ml of hexane) drop-wise under N₂ at -78°C. After 1 hour, MgBr•Et₂O (12.38g, 48mmol) was added and the reaction mixture was allowed to warm to -45°C for 1 hour before being treated with 2-Boc-Nlu-aldehyde (3.2g, 24mmol) in THF (20ml). The reaction mixture was stirred for 1 hour, quenched with saturated NH₄Cl, and extracted with ethyl acetate. The organic layer was washed with brine, dried with MgSO₄ and concentrated. The residue was purified by silica gel column chromatography to yield {1-[(5-ethyl-[1,3,4]oxadiazol-2-yl)-hydroxy-methyl]-propyl}-carbamic acid *tert*-butyl ester (2.13g). ¹ NMR (DMSO-*d*₆): 6.65, 6.52(1H, d, J=9.2Hz, J=9.2Hz, NH, diastereomer), 6.14, 5.95(1H, d, d, J=5.6Hz, J=5.6Hz, OH, diastereomer), 4.758, 4.467(1H, m, diastereomer), 3.7-3.55(1H, m), 2.8(2H, q), 1.33(12H, t), 1.25-1.21(2H, m), 0.82(3H, m). MS: 284.1 (M-1), 286 (M+1), 308(M+Na).

REFERENCE EXAMPLE 17

(a) (S)-2-Amino-1-benzooxazol-2-yl-butan-1-ol

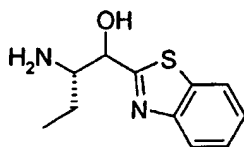


Step 1. Benzoxazole (600 mg, 5 mmol) in 20 ml THF was cooled to -5°C and isopropyl magnesium chloride (2M in THF, 2.5 ml, 5 mmol) was added. After stirring for 1 hour at -5°C, (S)-(1-formyl-propyl)-carbamic acid *tert*-butyl ester (561 mg, 3 mmol, Reference Example 18(a)), prepared as in reference 15, in 10 ml THF was added. The reaction was allowed to warm to room temperature with stirring for 2 hours. The reaction was quenched with saturated ammonium chloride solution, excess THF solvent removed. The residue was extracted with EtOAc, washed with brine, dried with anhydrous MgSO₄, filtered and concentrated. The crude residue was purified by chromatograph to yield 688 mg product (75%); LC-MS: 305.2 (M-1), 307.0 (M+1).

Step 2. (S)-[1-(Benzooxazol-2-yl-hydroxy-methyl)-propyl]-carbamic acid *tert*-butyl ester (275mg, 0.89mmol) and MeCl₂ (5ml) were mixed and TFA (1ml) was added at room temperature. After stirring for 1 hour, the solvent and excess TFA were removed under vacuum to produce 260mg of (S)-2-amino-1-benzooxazol-2-yl-butan-1-ol TFA salt.

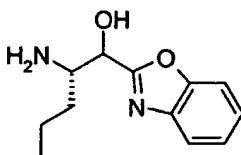
(b) (S)-2-Amino-1-benzothiazol-2-yl-butan-1-ol

-75-



By proceeding in a similar manner to Example 17(a) but using benzothiazole in Step 1 there was prepared (S)-2-amino-1-benzothiazol-2-yl-butan-1-ol TFA salt.

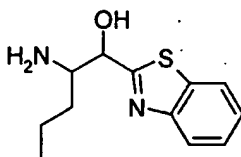
5 (c) (S)-2-amino-1-benzooxazol-2-yl-pentan-1-ol



By proceeding in a similar manner to Example 17(a) but using (S)-(1-formyl-butyl)-carbamic acid tert-butyl ester {561 mg, 3 mmol, Reference Example 18(b)} in Step 1 there was prepared (S)-2-amino-1-benzooxazol-2-yl-pentan-1-ol.

10

(d) 2-amino-1-benzothiazol-2-yl-pentan-1-ol



By proceeding in a similar manner to Example 17(a) but using benzothiazole and (S)-(1-formyl-butyl)-carbamic acid tert-butyl ester {561 mg, 3 mmol, Reference Example 18(b)} in Step 1 there was prepared 2-amino-1-benzothiazol-2-yl-pentan-1-ol.

15

REFERENCE EXAMPLE 18

20 (a) (S)-(1-Formyl-propyl)-carbamic acid tert-butyl ester

(S)-(+)-2-amino-1-butanol (50g, 561mmol) in 200ml of water and 200ml dioxane was cooled to 0°C and mixed with NaOH (26.9g, 673mmol) and di-t-butyl-dicarbonate (146.96 g, 673mmol). After the addition, the reaction was allowed to warm to room temperature. The reaction mixture was stirred for 2 hours. After removing the dioxane, the residue was extracted with EtOAc, then washed with brine and dried with anhydrous MgSO₄, filtered and concentrated. Without further purification, the crude product (120g) was used for next step reaction.

25

A solution of oxlyl chloride (40.39 g, 265mmol) in 700ml of MeCl₂ was stirred and cooled to -60°C. Dimethylsulfoxide (51.7 g, 663mmol) in 100 ml of MeCl₂ was added drop wise. After 10

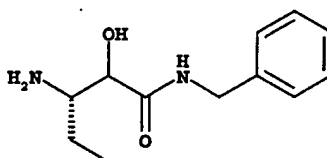
-76-

minutes a solution of (S)-2-boc-amino-1-butanol (50 g, 265 mmol) in 100ml of MeCl₂ was added drop wise at -70°C. The reaction mixture was allowed to warm to -40°C for 10 minutes and then cooled to -70°C again. A solution of triethylamine (74.9 g, 742mmol) in 100 ml of MeCl₂ was added. The reaction mixture was allowed to warm to room temperature over 2 hours. 100mls of saturated sodium dihydrogen phosphate was added, and then the organic layer was washed with brine and dried over MgSO₄. The solvent was removed to yield 45g of (S)-(1-formyl-propyl)-carbamic acid tert-butyl ester; H¹ NMR (DMSO-*d*₆): 9.4(1H, s), 7.29(1H, br.), 3.72(1H, m), 1.69(2H, m), 1.4-1.2(9H, s), 0.86(3H, t).

(b) By proceeding in a similar manner to Reference Example 18(a) but using (S)-(+)-2-amino-1-pentanol there was prepared (S)-(1-formyl-butyl)-carbamic acid tert-butyl ester.

REFERENCE EXAMPLE 19

(S)-3-Amino-2-hydroxy-pentanoic acid benzylamide



Step 1. (1S)-(2-Cyano-1-ethyl-2-hydroxyethyl)carbamic acid tert-butyl ester (10g, 46.7mmol) was dissolved in 1,4-dioxane (100mL). Anisole (5mL) was added and then concentrated HCl (100mL). The mixture was heated under reflux for 24 hours. The mixture was evaporated to dryness under vacuum and re-dissolved in 100mL water. The solution was washed with ether and then neutralized with saturated aqueous NaHCO₃. Di-tert-butyl dicarbonate (10g, 46mmol) was added with 1,4-dioxane (200mL), and the mixture was stirred at ambient temperature for 24 hours. The dioxane was removed under vacuum and the remaining aqueous solution was washed with ether. The solution was acidified with 1N HCl and extracted with ethyl acetate. The combined organic layers were washed with brine, dried with magnesium sulfate and evaporated to yield 3-tert-Butoxycarbonylamino-2-hydroxy-pentanoic acid (4.5g) as yellowish oil.

Step 2. 3-tert-Butoxycarbonylamino-2-hydroxy-pentanoic acid (300mg, 1.29mmol) was combined with EDC (400mg, 2.1mmol) and HOBt (400mg, 2.6mmol). A solution of benzylamine (0.22mL) and 4-methylmorpholine (0.5mL) in dichloromethyl (4mL) was added in one portion. The mixture was stirred at ambient temperature for 2 hours. After dilution with ethyl acetate (150mL), the solution was washed with 1N aqueous HCl, water, saturated aqueous NaHCO₃ solution and brine. The resultant mixture was dried with magnesium sulfate and evaporated under vacuum to yield (S)-3-amino-2-hydroxy-pentanoic acid benzylamide (380mg) as a white solid.

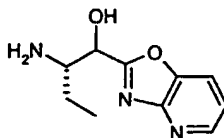
-77-

Step 3. (S)-3-Amino-2-hydroxy-pentanoic acid benzylamide was dissolved in a mixture of TFA/dichloromethyl (1:1; 6mL), stirred for 1 hour and evaporated to dryness. (3S)-3-Amino-2-hydroxy-pentanoic acid benzylamide was obtained as the TFA salt and used without further purification.

5

REFERENCE EXAMPLE 20

(S)-2-Amino-1-oxazolo[4,5-b]pyridin-2-yl-butan-1-ol



Step 1. A mixture of 2-amino-3-hydroxy pyridine (25g, 227mmol), triethylorthoformate (75ml) and p-toluenesulfonic acid (61mg) was heated at 140°C for 8 hours. Excess triethylorthoformate was removed under vacuum. The product was crystallized from ethyl acetate to yield 22.5g of pyridyloxazole; ^1H NMR (DMSO- δ): 9.26 (1H, s), 8.78 (1H, d), 8.45 (1H, d), 7.7(1H, dd); MS: 120.8 (M+1).

Step 2. Pyridyloxazole (600 mg, 5 mmol) in 30 ml THF was cooled to 0°C before the addition of isopropanyl magnesium chloride (2M in THF, 2.5 ml, 5 mmol). After stirring for 1 hour at 0°C, (S)-(1-formyl-propyl)-carbamic acid tert-butyl ester (573 mg, 3 mmol, Reference Example 18) in 20 ml THF was added. The ice bath was removed and the reaction allowed to warm to room temperature. The reaction mixture was stirred for 2 hours and quenched with saturated ammonium chloride solution. Excess THF was removed and the residue was extracted with EtOAc, washed with brine, dried with anhydrous MgSO_4 , filtered and concentrated. The crude residue was purified by chromatography to yield [1-(hydroxy-oxazolo[4,5-b]pyridin-2-yl-methyl)-propyl]-carbamic acid tert-butyl ester (383 mg) ^1H NMR (DMSO- δ): 8.42(1H, m), 8.18(1H, m), 7.3(1H, m), 6.8, 6.6(1H, dd, d, OH, diastereomeric), 6.3, 6.02(1H, d, d, NH, diastereomeric), 4.82, 4.5(1H, m, m, diastereomeric), 1.8-1.3(2H, m), 1.2, 1.05(9H, s, s, diastereomeric), 0.89(3H, m); MS: 306.2(M-1), 308.6(M+1).

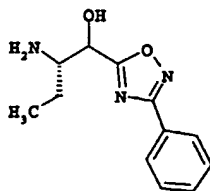
Step 3. To a stirred solution of the [1-(hydroxy-oxazolo[4,5-b]pyridin-2-yl-methyl)-propyl]-carbamic acid tert-butyl ester (12g, 100mmol) in THF (300ml) was added n-BuLi (1.6M solution in 62.5ml of hexane) drop wise under N_2 at -78°C. After 1 hour, $\text{MgBr}\cdot\text{Et}_2\text{O}$ (25.8g, 100mmol) was added and the reaction mixture was allowed to warm to -45°C for 1 hour before being treated with 2-boc-amino-butyl-aldehyde (11.46g, 60mmol) in THF (50ml). The reaction mixture was stirred for 1 hour, quenched with saturated NH_4Cl , and extracted with ethyl acetate. The organic layer was washed with brine, dried with MgSO_4 and concentrated. The residue was purified by silica gel column chromatography to yield 2-boc-amino-1-(5-pyridyloxazole-2-yl)-1-butanol (14.1g).

-78-

Step 4. 2-Boc-amino-1-(5-pyridyloxazole-2-yl)-1-butanol (311mg, 1mmol) and MeCl₂ (5ml) were mixed and TFA (1ml) was added at room temperature. After stirring for 1 hour, the solvent and excess TFA were removed under vacuum to produce 355mg of 2-amino-1-oxazol[4,5-b]pyridin-2-yl-butan-1-ol TFA salt.

REFERENCE EXAMPLE 21

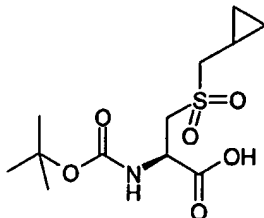
(S)-2-Amino-1-(3-phenyl-[1,2,4]oxadiazol-5-yl)-butan-1-ol



3-*tert*-Butoxycarbonylamino-2-hydroxy-pentanoic acid (500mg, 2.14mmol) was combined with EDC (600mg, 3.14mmol), HOBt (600mg, 3.92mmol), and N-hydroxy-benzamidine (292mg, 2.14mmol). Dichloromethyl (10mL) was added and then 4-methylmorpholine (1mL). The mixture was stirred at ambient temperature for 16 hours. After dilution with ethyl acetate (200mL), the solution was washed with water (30mL), saturated aqueous NaHCO₃ solution and brine, dried with MgSO₄ and evaporated under vacuum. The crude product was dissolved in pyridine (10mL) and heated at 80°C for 15 hours. The pyridine was evaporated under vacuum and the residue was purified by flash chromatography on silica gel (eluent: ethyl acetate) to yield 290mg (0.83mmol). The oxadiazole (145mg, 0.41mmol) was dissolved in CH₂Cl₂ (4mL) and TFA (4mL) was added. After stirring for 1 hour, the mixture was evaporated to dryness to yield (S)-2-Amino-1-(3-phenyl-[1,2,4]oxadiazol-5-yl)-butan-1-ol.

REFERENCE EXAMPLE 22

(R)-2-*tert*-butoxycarbonylamino-3-cyclopropylmethanesulfonyl-propionic acid



Step 1. Sodium hydroxide (2.16g, 54mmol) was dissolved in 27ml water and the solution added to a suspension of (R)-2-*tert*-butoxycarbonylamino-3-mercapto-propionic acid (8.2g, 37mmol) in 54ml methanol. After a clear solution had formed bromomethyl-cyclopropane (5g, 37mmol) was added and the resulting reaction mixture stirred for three days. Methanol was removed under reduced pressure.

-79-

The residue was treated with 200ml 1M hydrochloric acid and then extracted three times with 200ml of dichloromethane. The combined organic phases were washed with brine and dried with magnesium sulfate. The solvent was evaporated under reduced pressure to give 2-tert-butoxycarbonylamino-3-cyclopropylmethylsulfanyl-propionic acid (7.94g).

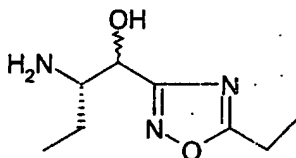
5

Step 2. Sodium hydroxide (2.32g, 58mmol) was dissolved in 75ml water. 2-tert-butoxycarbonylamino-3-cyclopropylmethylsulfanyl-propionic acid (7.94g, 29mmol) was added. A solution of Oxone™ in 100ml water was added slowly. The pH was adjusted to 3 by addition of sodium bicarbonate and the reaction mixture stirred for 30 minutes. It was extracted three times with 100ml ethyl acetate. The combined organic phases were washed with 100ml brine and dried with magnesium sulfate. The solvent was removed to yield (R)-2-tert-butoxycarbonylamino-3-cyclopropylmethanesulfonyl-propionic acid (4.64g, 15mmol, 31%).

10

REFERENCE EXAMPLE 23

15 (S)-2-Amino-1-(5-ethyl-1,2,4-oxadiazol-3-yl)-butan-1-ol trifluoro-acetic acid salt



Step 1. A solution of (2-Cyano-1-ethyl-2-hydroxy-ethyl)-carbamic acid tert-butyl ester (1, 9.53g, 44 mmol) in methanol (80 ml) was cooled to 0°C and treated successively with hydroxylamine hydrochloride (3.05, 44 mmol) in methanol (80 ml) and 25% sodium methoxide solution in methanol (10.2 ml). Stirred at 0°C for 5 min., cold bath removed and the reaction mixture stirred at room temperature for 5hr. Methanol evaporated off under reduced pressure, crude partitioned between ethyl acetate and water. Organic layer separated, dried (MgSO₄) and evaporated under reduced pressure to give yellow oil. Purified by mpic eluting with a mixture of ethyl acetate – heptane to give {(S)-1-[Hydroxy-(N!-hydroxycarbamimidoyl)-methyl]-propyl}-carbamic acid tert-butyl ester as white solid (3.5 g). MS: M(H⁺) 248.

25

Step 2. A mixture of {(S)-1-[Hydroxy-(N!-hydroxycarbamimidoyl)-methyl]-propyl}-carbamic acid tert-butyl ester (525 mg, 2.16 mmol), propionic anhydride (0.3 ml, 2.37 mmol) in dioxane (5ml) was heated at 150°C in a microwave (Smith Creator, S00219) for 35min. Crude evaporated under reduced pressure and purified by flash column chromatography to give {(S)-1-[(5-Ethyl-1,2,4-oxadiazol-3-yl)-hydroxy-methyl]-propyl}-carbamic acid tert-butyl ester as yellow solid (0.8g, 67%).

30

¹H NMR (CDCl₃): 4.88-4.80 (2H, m), 4.01-3.84 (1H, 2 broad m), 3.64-3.45 (1H, 2 bs), 2.95-2.86 (2H, dq, J=4.2Hz, 7.6Hz), 1.73-1.62 (1H, m), 1.6-1.32 (13H, m), 1.02-0.94 (3H, q, J=7.5Hz). MS:

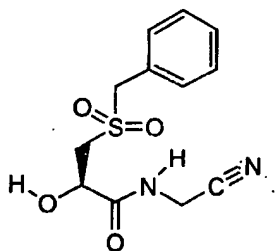
-80-

304(M+1)

- Step 3. {(S)-1-[(5-Ethyl-1,2,4-oxadiazol-3-yl)-hydroxy-methyl]-propyl}-carbamic acid tert-butyl ester (214 mg, 0.75 mmol) in dichloromethane (3 ml)) was treated with trifluoro acetic acid at room temperature for 3h. Solvent evaporated under reduced pressure to give (S)-2-Amino-1-(5-ethyl-1,2,4-oxadiazol-3-yl)-butan-1-ol trifluoro-acetic acid salt as brown oil (0.3 g). ¹H NMR (CDCl₃): 7.9-7.4(3H, 2bs), 5.07 & 5.24 (1H, 2 x d, J=3.5Hz & 5.5Hz), 3.8-3.6 (1H, 2 bs), 2.96-2.87 (2H, dq, J=2.4Hz, 7.5Hz), 1.8-1.4 (2H, m), 1.40-1.34 (3H, dt, J=1.4Hz, 7.5Hz), 1.06-0.98 (3H, dt, J=7.5Hz, 10.5Hz).
- MS: 186(M+1)

EXAMPLE 1

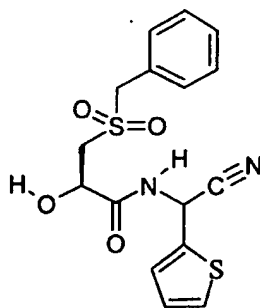
- (a) (R)-N-cyanomethyl-2-hydroxy-3-phenylmethanesulfonyl-propionamide, (Compound 4)



- DMF (5mL) was added to a mixture of 2-hydroxy-3-phenylmethanesulfonyl-propionic acid [200mg, 0.82mmol, Reference Example 1(b)], EDC (300mg, 1.57mmol), HOBt (300mg, 1.96mmol) and aminoacetonitrile hydrochloride (200mg, 2.1mmol). 4-Methylmorpholine (0.5mL) was added and the mixture was stirred at ambient temperature for 2 hours. The mixture was diluted with ethyl acetate (200mL), washed with 1N HCl, brine, saturated aqueous NaHCO₃ solution, and brine, dried with MgSO₄ and evaporated under vacuum. (R)-N-cyanomethyl-2-hydroxy-3-phenylmethanesulfonyl-propionamide was crystallized from ethyl acetate/hexane to yield 154mg (0.55mmol); ¹H NMR: (DMSO) 8.89-8.77 (m, 1H), 7.46-7.37 (m, 5H), 6.71-6.62 (m, 1H), 4.60-4.45 (m, 3H), 4.17-4.08 (m, 2H), 3.39-3.28 (m, 2H). MS: (M⁺+1) 283.

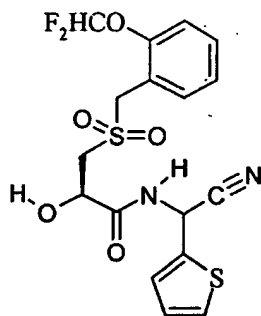
- (b) (R)-N-(1-cyano-1-thiophen-2-yl-methyl)-2-hydroxy-3-phenylmethanesulfonyl-propionamide, (Compound 7);

-81-



By proceeding in a manner similar to Example 1(a) above but using (R)-2-hydroxy-3-phenylmethanesulfonyl-propionic acid [Reference Example 1(b)] and DL- α -amino-2-thiopheneacetic acid there was prepared (R)-N-(1-cyano-1-thiophen-2-yl-methyl)-2-hydroxy-3-phenylmethanesulfonyl-propionamide. ¹H NMR (DMSO): 9.55(d, J=6.5Hz, 1H), 7.58(d, J=5.21Hz, 1H), 7.42-7.39(m, 5H), 7.23(m, 1H), 7.05(dd, J=3.51Hz, J=5.21Hz, 1H), 6.58(dd, J=3.45Hz, J=6.66Hz, 1H), 6.41(s, 1H), 4.59-4.50(m, 3H), 3.29(s, 2H); MS: 362.6(M⁻), 365(M⁺).

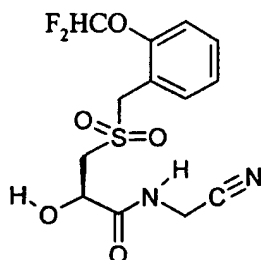
- (c) (R)-N-(1-cyano-1-thiophen-2-yl-methyl)-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionamide, (Compound 8)



- By proceeding in a manner similar to Example 1(a) above but using (R)-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionic acid [Reference Example 1(a)] and DL- α -amino-2-thiopheneacetic acid there was prepared (R)-N-(1-cyano-1-thiophen-2-yl-methyl)-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionamide. ¹H NMR (CD₃Cl): δ 7.6-7.2(m, 7H), 7.01(t, J=73.6Hz, 1H), 6.62(s, 1H), 6.21(d, J=8.15, 1H), 4.71-4.67(m, 1H), 4.46(s, 2H), 3.68(s, 2H), 3.22-3.18(m, 1H); MS: 428.6(M-1), 453(M+23).

- (d) (R)-N-cyanomethyl-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionamide, (Compound 17)

-82-

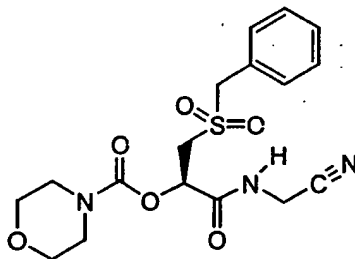


By proceeding in a manner similar to Example 1(a) above but using (R)-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionic acid [Reference Example 1(a)] there was prepared (R)-N-cyanomethyl-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionamide.

- 5 ¹HNMR (DMSO): 8.81(t, J=5.67Hz, 1H), 7.55-7.4(m, 2H), 7.35-7.2(m, 2H), 7.13(t, J=73.68Hz, 1H), 6.62(d, J=6.67Hz, 1H), 4.58(s, 2H), 4.52-4.45(m, 1H), 4.12(d, J=5.94Hz, 2H), 3.45-3.4(m, 2H). MS: 347.4(M-1), 371(M+23).

EXAMPLE 2

- 10 Morpholine-4-carboxylic acid (R)-1-(cyanomethyl-carbamoyl)-2-phenylmethanesulfonyl-ethyl ester, (Compound 6);

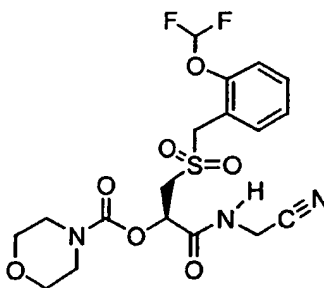


- Phosgene solution (0.77mL, 1.93M in toluene) was added to CH₂Cl₂ (5mL) and cooled to 0°C under nitrogen. Quinoline (0.12mL, 1.0mmol) was added followed by (R)-N-cyanomethyl-2-hydroxy-3-phenylmethanesulfonyl-propionamide [100mg, 0.354mmol, Example 1(a)]. The mixture was stirred at ambient temperature for 3 hours. Morpholine (1mmol) was added and stirring was continued for 3 hours. The mixture was diluted with ethyl acetate (200mL) and washed sequentially with 1N HCl, brine, saturated aqueous NaHCO₃ solution and brine. The product was dried with MgSO₄ and evaporated under vacuum and crystallized from an ethyl acetate/hexane solution to yield morpholine-4-carboxylic acid (R)-1-(cyanomethyl-carbamoyl)-2-phenylmethanesulfonyl-ethyl ester. (85mg; 0.215mmol); ¹H NMR: (DMSO) 8.99-8.88 (m, 1H), 7.46-7.37 (m, 5H), 5.42-5.32 (m, 1H), 4.60-4.45 (m, 2H), 4.20-4.13 (m, 2H), 3.70-3.28 (m, 10H). MS: (M⁺+1) 396.

EXAMPLE 3

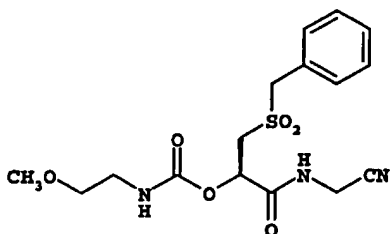
- 25 (a) Morpholine-4-carboxylic acid (R)-1-(cyanomethyl-carbamoyl)-2-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-ethyl ester, (Compound 31)

-83-



(*R*)-*N*-cyanomethyl-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionamide [100mg, 0.287mmol, Example 1(d)], was dissolved in CH₂Cl₂ (2mL). Pyridine (32.4μL, 0.4mmol) and then trichloromethylchloroformate (36.2μL, 0.3mmol) were added. The mixture was stirred at ambient temperature for 3 hours. Morpholine (0.5mL) was added and stirring was continued for 3 hours. The mixture was diluted with ethyl acetate (200mL), washed with 1N HCl, brine, saturated aqueous NaHCO₃ solution and brine. The product was dried with MgSO₄, evaporated under vacuum and crystallized from a solution of ethyl acetate/hexane to yield morpholine-4-carboxylic acid (*R*)-1-(cyanomethyl-carbamoyl)-2-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-ethyl ester (60mg; 0.130mmol); ¹H NMR: (DMSO) δ 8.95 (t, J=5.2Hz, 1H), 7.51-7.44 (m, 2H), 7.32-7.22 (m, 2H), 7.14 (t, J_{HF}=73Hz, 1H), 5.39-5.35 (m, 1H), 4.67-4.53 (m, 2H), 4.19-4.15 (m, 2H), 3.83-3.28 (m, 10H); MS: (M⁺+1) 462.

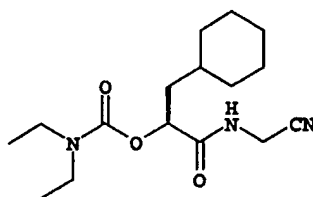
(b) (*R*)-(2-Methoxy-ethyl)-carbamic acid 1-(cyanomethyl-carbamoyl)-2-phenylmethanesulfonyl-ethyl ester



By proceeding in a manner similar to Example 3(a) above but using (*R*)-*N*-cyanomethyl-2-hydroxy-3-phenylmethanesulfonyl-propionamide [Example 1(a)] and 2-methoxyethylamine there was prepared (*R*)-(2-Methoxy-ethyl)-carbamic acid 1-(cyanomethyl-carbamoyl)-2-phenylmethanesulfonyl-ethyl ester. ¹H NMR: (DMSO) 8.91 (t, J=5.6Hz, 1H), 7.64 (t, J=5.6Hz, 1H), 7.40-7.32 (m, 5H), 5.30-5.25 (m, 1H), 4.59-4.50 (m, 2H), 4.17-4.13 (m, 2H), 3.58 (dd, J=9.2Hz, J=14.8Hz, 1H), 3.43 (d, 14.8Hz, 1H), 3.33 (s, 3H), 3.38-3.12 (m, 4H). MS: (M+H)⁺ 384.

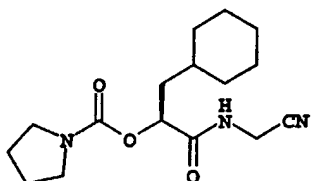
(c) (*S*)-Diethyl-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester

-84-



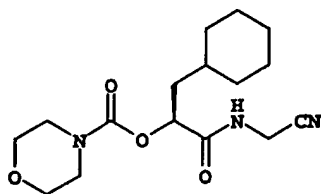
By proceeding in a manner similar to Example 3(a) above but using (*R*)-*N*-cyanomethyl-3-cyclohexyl-2-hydroxy-propionamide and diethylamine there was prepared (*S*)-Diethyl-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester. ¹H NMR: (DMSO) 8.62 (t, J=5.6Hz, 1H), 4.87-4.82 (m, 1H), 4.12 (d, J=5.6, 2H), 3.42-3.10 (m, 4H), 1.72-0.82 (m, 19H). MS: (M+H)⁺ 310.

(d) (*S*)-Pyrrolidine-1-carboxylic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester



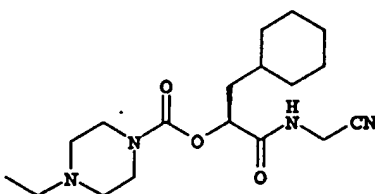
By proceeding in a manner similar to Example 3(a) above but using (*R*)-*N*-cyanomethyl-3-cyclohexyl-2-hydroxy-propionamide and pyrrolidine there was prepared (*S*)-Pyrrolidine-1-carboxylic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester. ¹H NMR: (DMSO) 8.59 (t, J=4.8Hz, 1H), 4.86-4.81 (m, 1H), 4.11 (d, J=4.8, 2H), 3.48-3.19 (m, 4H), 1.87-0.82 (m, 17H). MS: (M+H)⁺ 308.

(e) (*S*)-Morpholine-4-carboxylic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester



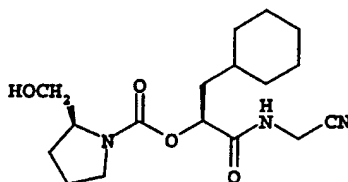
By proceeding in a manner similar to Example 3(a) above but using (*R*)-*N*-cyanomethyl-3-cyclohexyl-2-hydroxy-propionamide and morpholine there was prepared (*S*)-Morpholine-4-carboxylic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester. ¹H NMR: (DMSO) 8.66 (t, J=5.2Hz, 1H), 4.88-4.83 (m, 1H), 4.13 (d, J=4.8, 2H), 3.60-3.26 (m, 8H), 1.71-0.82 (m, 13H). MS: (M+H)⁺ 324.

(f) (*S*)-4-Ethyl-piperazine-1-carboxylic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester



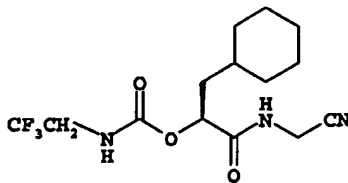
By proceeding in a manner similar to Example 3(a) above but using (*R*)-*N*-cyanomethyl-3-cyclohexyl-2-hydroxy-propionamide and 4-ethylpiperazine there was prepared (S)-4-Ethyl-piperazine-1-carboxylic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester. LC-MS: elution time = 2.08min. 349.2(M-1), 351.3(M+1). (MS: API 150EX. LC: HP Agilent 1100 Series. Column: Phenomenex, 5u ODS3 100A 100X3mm.; Flow Rate: 2ml/min. Two solvent gradient: Solvent A, 99% water, 1% acetonitrile, 0.1% AcOH. Solvent B, 99% acetonitrile, 1% water, 0.1% AcOH. Gradient from 100% A, 0% B to 0% A, 100% B from t = 0 to t = 6min. Then gradient back to 100% A, 0% B from t = 7 to t = 15 min.).

- 10 (g) (S)-2-Hydroxymethyl-pyrrolidine-1-carboxylic acid (S)-1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester



- By proceeding in a manner similar to Example 3(a) above but using (*R*)-*N*-cyanomethyl-3-cyclohexyl-2-hydroxy-propionamide and (S)-2-hydroxymethyl-pyrrolidine there was prepared (S)-2-Hydroxymethyl-pyrrolidine-1-carboxylic acid (S)-1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester. LC-MS: elution time = 3.73min. 336.5(M-1), 338.2(M+1). (MS: API 150EX. LC: HP Agilent 1100 Series. Column: Phenomenex, 5u ODS3 100A 100X3mm.; Flow Rate: 2ml/min. Two solvent gradient: Solvent A, 99% water, 1% acetonitrile, 0.1% AcOH. Solvent B, 99% acetonitrile, 1% water, 0.1% AcOH. Gradient from 100% A, 0% B to 0% A, 100% B from t = 0 to t = 6min. Then gradient back to 100% A, 0% B from t = 7 to t = 15 min.)

- (h) (S)-(2,2,2-Trifluoro-ethyl)-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester

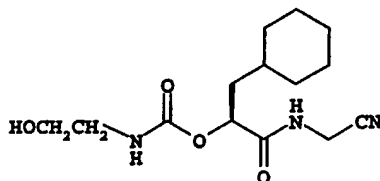


- By proceeding in a manner similar to Example 3(a) above but using (*R*)-*N*-cyanomethyl-3-cyclohexyl-2-hydroxy-propionamide and 2,2,2-trifluoroethylamine there was prepared (S)-(2,2,2-Trifluoro-ethyl)-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester. LC-MS: elution time = 4.07min. 334.1(M-1), 336.2(M+1). (MS: API 150EX. LC: HP Agilent 1100 Series. Column: Phenomenex, 5u ODS3 100A 100X3mm.; Flow Rate: 2ml/min. Two solvent gradient: Solvent A, 99% water, 1% acetonitrile, 0.1% AcOH. Solvent B, 99% acetonitrile, 1% water, 0.1% AcOH. Gradient from 100% A,

-86-

0% B to 0% A, 100% B from $t = 0$ to $t = 6$ min. Then gradient back to 100% A, 0% B from $t = 7$ to $t = 15$ min.)

(i) (S)-(2-Hydroxyethyl)-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester

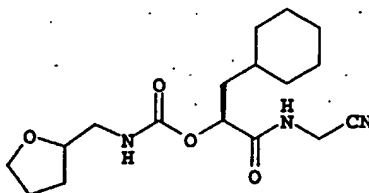


5

By proceeding in a manner similar to Example 3(a) above but using (*R*)-*N*-cyanomethyl-3-cyclohexyl-2-hydroxy-propionamide and 2-hydroxyethylamine there was prepared (S)-(2-Hydroxyethyl)-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester. ^1H NMR: (DMSO) 8.65 (t, $J=5.2\text{Hz}$, 1H), 7.16 (t, $J=5.2\text{Hz}$, 1H), 4.85-4.80 (m, 1H), 4.62 (t, $J=5.6\text{Hz}$, 1H), 4.12 (d, $J=5.6\text{Hz}$, 2H), 3.45-3.33 (m, 2H), 3.12-2.96 (m, 2H), 1.74-0.80 (m, 13H). MS: $(\text{M}+\text{H})^+$ 298.

10

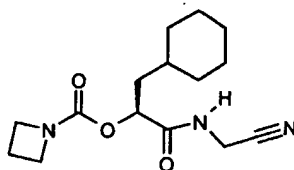
(j) (Tetrahydrofuran-2-ylmethyl)-carbamic acid (S)-1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester



15 By proceeding in a manner similar to Example 3(a) above but using (*R*)-*N*-cyanomethyl-3-cyclohexyl-2-hydroxy-propionamide and tetrahydrofurfurylamine there was prepared (tetrahydrofuran-2-ylmethyl)-carbamic acid (S)-1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester as a 1:1 mixture of diastereomers, ^1H NMR: (DMSO) 8.66 (t, $J=5.2\text{Hz}$, 1H), 7.28 (t, $J=5.2\text{Hz}$, 1H), 4.86-4.81 (m, 1H), 4.12 (d, $J=5.2\text{Hz}$, 2H), 3.83-3.54 (m, 3H), 3.09-2.92 (m, 2H), 1.89-0.80 (m, 17H). MS: $(\text{M}+\text{H})^+$ 338.

20

(k) (S)-Azetidine-1-carboxylic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester



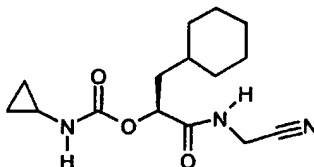
By proceeding in a manner similar to Example 3(a) above but using (*R*)-*N*-cyanomethyl-3-cyclohexyl-2-hydroxy-propionamide and azetidine there was prepared (S)-azetidine-1-carboxylic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester. ^1H NMR: (DMSO) 8.59 (t, $J=5.2\text{Hz}$, 1H), 4.82-

25

-87-

4.77 (m, 1H), 4.11 (d, J=5.2Hz, 2H), 4.13-3.81 (m, 4H), 2.18 (quint, J=7.6Hz, 2H), 1.71-0.80 (m, 13H). MS: (M+H)⁺ 294.

(l) (S)-Cyclopropyl-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester

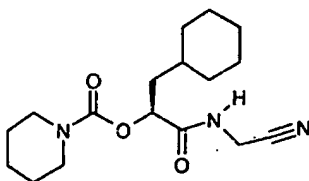


5

By proceeding in a manner similar to Example 3(a) above but using (*R*)-*N*-cyanomethyl-3-cyclohexyl-2-hydroxy-propionamide and cyclopropylamine there was prepared (*S*)-cyclopropyl-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester. ¹H NMR: (DMSO) 8.64 (t, J=5.2Hz, 1H), 7.44 (br, 1H), 4.83-4.78 (m, 1H), 4.11 (d, J=5.2Hz, 2H), 2.50-2.40 (m, 1H), 1.72-0.78 (m, 13H), 0.58-0.30 (m, 4H). MS: (M+H)⁺ 294.

10

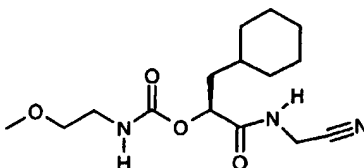
(m) (S)-Piperidine-1-carboxylic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester



15

By proceeding in a manner similar to Example 3(a) above but using (*R*)-*N*-cyanomethyl-3-cyclohexyl-2-hydroxy-propionamide and piperidine there was prepared (*S*)-piperidine-1-carboxylic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester. ¹H NMR: (DMSO) 8.63 (t, J=5.2Hz, 1H), 4.86-4.81 (m, 1H), 4.11 (d, J=5.6Hz, 2H), 3.48-3.20 (m, 4H), 1.70-0.82 (m, 19H). MS: (M+H)⁺ 322.

(n) (S)-(2-Methoxy-ethyl)-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester



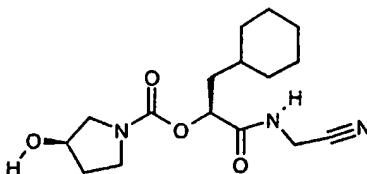
20

By proceeding in a manner similar to Example 3(a) above but using (*R*)-*N*-cyanomethyl-3-cyclohexyl-2-hydroxy-propionamide and 2-methoxyethylamine there was prepared (*S*)-(2-methoxy-ethyl)-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester. ¹H NMR: (DMSO) 8.66 (t, J=5.6Hz, 1H), 7.27 (t, J=5.6Hz, 1H), 4.85-4.80 (m, 1H), 4.12 (d, J=5.6Hz, 2H), 3.40-3.03 (m, 4H), 3.32 (s, 3H), 1.74-0.80 (m, 13H). MS: (M+H)⁺ 312.

25

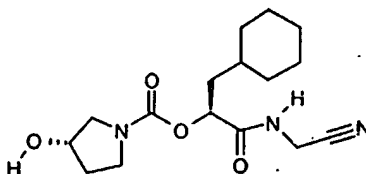
-88-

- (o) (R)-3-Hydroxy-pyrrolidine-1-carboxylic acid (S)-1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester



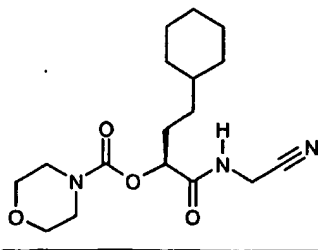
- By proceeding in a manner similar to Example 3(a) above but using (R)-N-cyanomethyl-3-cyclohexyl-2-hydroxy-propionamide and (R)-3-hydroxy-pyrrolidine there was prepared (R)-3-hydroxy-pyrrolidine-1-carboxylic acid (S)-1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester. ¹H NMR: (DMSO) 8.64-8.56 (m, 1H), 4.98-4.80 (m, 2H), 4.29-4.20 (m, 1H), 4.11 (d, J=5.2Hz, 2H), 3.57-3.12 (m, 4H), 1.91-0.82 (m, 15H). MS: (M+H)⁺ 324.

- (p) (S)-3-Hydroxy-pyrrolidine-1-carboxylic acid (S)-1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester



- By proceeding in a manner similar to Example 3(a) above but using (R)-N-cyanomethyl-3-cyclohexyl-2-hydroxy-propionamide and (S)-3-hydroxy-pyrrolidine there was prepared (S)-3-hydroxy-pyrrolidine-1-carboxylic acid (S)-1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester. ¹H NMR: (DMSO) 8.63-8.55 (m, 1H), 4.98-4.90 (m, 1H), 4.85-4.80 (m, 1H), 4.28-4.19 (m, 1H), 4.13-4.09 (m, 2H), 3.54-3.09 (m, 4H), 1.93-0.81 (m, 15H). MS: (M+H)⁺ 324.

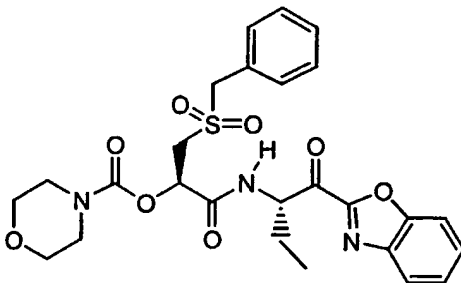
- (q) (S)-Morpholine-4-carboxylic acid 1-(cyanomethyl-carbamoyl)-3-cyclohexyl-propyl ester



- By proceeding in a manner similar to Example 3(a) above but using (R)-N-cyanomethyl-3-cyclohexyl-2-hydroxy-propionamide and morpholine there was prepared (S)-morpholine-4-carboxylic acid 1-(cyanomethyl-carbamoyl)-3-cyclohexyl-propyl ester. ¹H NMR: (DMSO) 8.61 (t, J=5.6Hz, 1H), 4.79 (t, J=5.6Hz, 1H), 4.13 (d, J=5.2Hz, 2H), 3.59-3.26 (m, 8H), 1.73-1.55 (m, 7H), 1.23-1.06 (m, 6H), 0.88-0.76 (m, 2H). MS: (M+H)⁺ 338.

EXAMPLE 4

- (a) Morpholine-4-carboxylic acid (R)-1-[(S)-1-(1-benzooxazol-2-yl-methanoyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester, (Compound 11)



5

Step 1. (R)-2-Hydroxy-3-phenylmethanesulfonyl-propionic acid {2g, 8.19mmol, Reference Example 1(b)} was dissolved in CH_2Cl_2 (20mL). 4-Methylmorpholine (3.8mL) and then chloromethyl methyl ether (1.52mL, 20mmol) were added. After stirring at ambient temperature for 30 minutes, the reaction was quenched with water (50mL) and extracted with ethyl acetate. The combined organic layers were washed with saturated aqueous NaHCO_3 solution and brine. The product was dried with MgSO_4 , evaporated under vacuum and crystallized from ethyl acetate/hexane to yield 2-hydroxy-3-phenylmethanesulfonyl-propionic acid methoxymethyl ester (1.2g; 4.16mmol).

10

Step 2. Phosgene solution (2.07mL, 1.93M in toluene) was added to CH_2Cl_2 (10mL) and cooled to 0°C under nitrogen. Quinoline (0.95mL) was added followed by 2-hydroxy-3-phenylmethanesulfonyl-propionic acid methoxymethyl ester (250mg, 0.87mmol). The mixture was stirred at ambient temperature for 3 hours. Morpholine (0.35mL, 4mmol) was added and stirring was continued for 3 hours. The mixture was diluted with ethyl acetate (200mL), washed sequentially with 1N HCl, brine, saturated aqueous NaHCO_3 solution and brine. The crude product was dried with MgSO_4 , evaporated under vacuum and dissolved in 1,4-dioxane (20mL). 1N HCl (10mL) was added and the mixture was stirred at ambient temperature for 3 hours. Dioxane was evaporated under vacuum and the product was extracted with ethyl acetate. The combined ethyl acetate layers were washed with saturated aqueous NaHCO_3 solution (3x20mL). The NaHCO_3 solution was acidified with 6N HCl and extracted with ethyl acetate. The combined ethyl acetate layers were washed with brine, dried with MgSO_4 and evaporated under vacuum to give (R)-morpholine-4-carboxylic acid 1-carboxy-2-phenylmethanesulfonyl-ethyl ester.

20

25

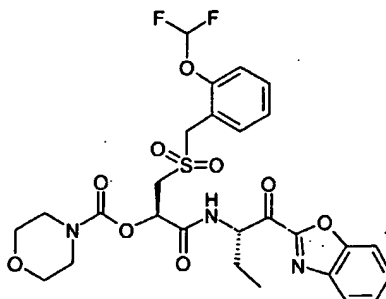
Step 3. (R)-Morpholine-4-carboxylic acid 1-carboxy-2-phenylmethanesulfonyl-ethyl ester was combined with EDC (250mg, 1.3mmol), HOBt (250mg, 1.6mmol), and (2S)-2-amino-1-benzooxazol-2-yl-butan-1-ol {250mg, 1.2mmol, Reference Example 17(a)}. Dichloromethane (4mL) was added and then 4-methylmorpholine (0.5mL). The mixture was stirred at ambient temperature for 2 hours. After dilution with ethyl acetate (150mL), the solution was washed with 1N aqueous HCl, water, saturated aqueous NaHCO_3 solution and brine, dried with MgSO_4 and evaporated under vacuum. The crude product was dissolved in dry dichloromethane (10mL) and

30

-90-

Dess Martin Periodinane (500mg, 1.2mmol) was added. After stirring at ambient temperature for 1 hour, the mixture was diluted with ethyl acetate (150mL) and treated with 0.26M Na₂S₂O₃ solution in saturated aqueous NaHCO₃. The organic phase was washed with saturated aqueous NaHCO₃ and brine, dried with MgSO₄ and evaporated. The product was crystallized from ethyl acetate/hexane to yield morpholine-4-carboxylic acid (R)-1-[(S)-1-(1-benzooxazol-2-yl-methanoyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester (190mg; 0.35mmol); ¹H NMR: (DMSO) 8.95 (d, J=6.6Hz, 1H), 8.01 (d, J=7.9Hz, 1H), 7.90 (d, J=7.9Hz, 1H), 7.65 (t, J=7.5Hz, 1H), 7.55 (t, J=7.9Hz, 1H), 7.40-7.34 (m, 5H), 5.44-5.35 (m 1H), 5.26-5.16 (m, 1H), 4.60 (d, J=13.6Hz, 1H), 4.47 (d, J=13.6Hz, 1H), 3.71-3.28 (m, 10H), 2.10-1.94 (m, 1H), 1.81-1.65 (m, 1H), 0.98 (t, J=7.2Hz, 3H); MS: (M⁺+1) 544.

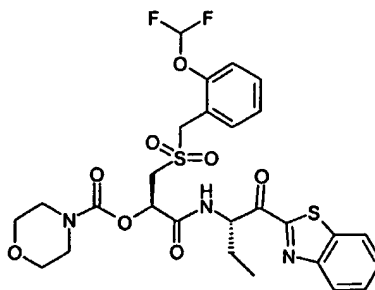
(b) Morpholine-4-carboxylic acid (R)-1-[(S)-1-(1-benzooxazol-2-yl-methanoyl)-propylcarbamoyl]-2-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-ethyl ester,
(Compound 14)



By proceeding in a manner similar to Example 4(a) above but using (R)-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionic acid {Reference Example 1(a)} in step 1 there was prepared morpholine-4-carboxylic acid (R)-1-[(S)-1-(1-benzooxazol-2-yl-methanoyl)-propylcarbamoyl]-2-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-ethyl ester ¹H NMR: (DMSO) 8.95 (d, J=6.4Hz, 1H), 8.01 (d, J=7.9Hz, 1H), 7.90 (d, J=8.4Hz, 1H), 7.65 (t, J=7.4Hz, 1H), 7.54 (t, J=7.5Hz, 1H), 7.52-7.43 (m, 2H), 7.31-7.21 (m, 2H), 7.11 (t, J_{H,F}=73Hz, 1H), 5.43-5.37 (m 1H), 5.27-5.17 (m, 1H), 4.63 (d, J=13.5Hz, 1H), 4.54 (d, J=13.5Hz, 1H), 3.88-3.28 (m, 10H), 2.10-1.94 (m, 1H), 1.81-1.65 (m, 1H), 0.98 (t, J=7.6Hz, 3H); MS: (M⁺+1) 610.

(c) morpholine-4-carboxylic acid (R)-1-[(S)-1-(1-benzothiazol-2-yl-methanoyl)-propylcarbamoyl]-2-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-ethyl ester,
(Compound 15).

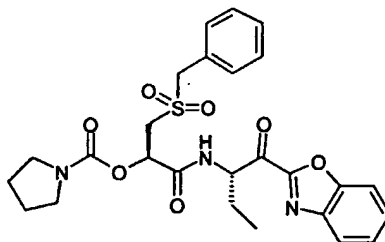
-91-



By proceeding in a manner similar to Example 4(a) above but using (R)-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionic acid {Reference Example 1(a)} in step 1 and (2S)-2-amino-1-benzothiazol-2-yl-butan-1-ol {Reference Example 17(b)} in step 3 there was prepared
 5 morpholine-4-carboxylic acid (R)-1-[(S)-1-(1-benzothiazol-2-yl-methanoyl)-propylcarbamoyl]-2-[(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-ethyl ester. ¹H NMR: (DMSO) 8.93 (d, J=6.4Hz, 1H), 8.30-8.24 (m, 2H), 7.72-7.62 (m, 2H), 7.51-7.44 (m, 2H), 7.32-7.22 (m, 2H), 7.12 (t, J_{H,F}=73Hz, 1H), 5.49-5.35 (m 2H), 4.64 (d, J=13.5Hz, 1H), 4.55 (d, J=13.5Hz, 1H), 3.91-3.28 (m, 10H), 2.08-1.94 (m, 1H), 1.84-1.68 (m, 1H), 0.99 (t, J=7.6Hz, 3H). MS: (M⁺+1) 626.

10

(d) Pyrrolidine-1-carboxylic acid (R)-1-[(S)-1-(1-benzooxazol-2-yl-methanoyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester, (Compound 19).

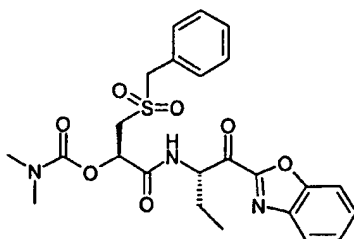


By proceeding in a manner similar to Example 4(a) above but using pyrrolidine in step 2 there was
 15 prepared pyrrolidine-1-carboxylic acid (R)-1-[(S)-1-(1-benzooxazol-2-yl-methanoyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester. ¹H NMR: (DMSO) 8.90 (d, J=6.4Hz, 1H), 7.99 (d, J=7.9Hz, 1H), 7.89 (d, J=8.4Hz, 1H), 7.65 (t, J=7.4Hz, 1H), 7.54 (t, J=7.5Hz, 1H), 7.40-7.33 (m, 5H), 5.41-5.33 (m 1H), 5.26-5.15 (m, 1H), 4.59 (d, J=13.5Hz, 1H), 4.47 (d, J=13.5Hz, 1H), 3.66-3.17 (m, 6H), 2.10-1.64 (m, 6H), 0.97 (t, J=7.2Hz, 3H); MS: (M⁺+1) 528.

20

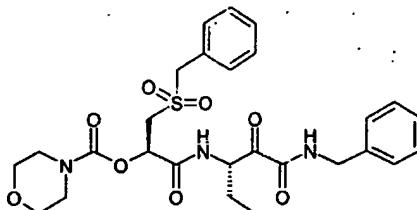
(e) Dimethyl-carbamic acid (R)-1-[(S)-1-(1-benzooxazol-2-yl-methanoyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester, (Compound 20).

-92-



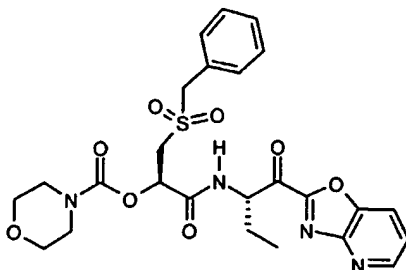
By proceeding in a manner similar to Example 4(a) above but using dimethylamine in step 2 there was prepared dimethyl-carbamic acid (R)-1-[(S)-1-(1-benzooxazol-2-yl-methanoyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester. ¹H NMR: (DMSO) 8.91 (d, J=6.4Hz, 1H), 7.99 (d, J=7.9Hz, 1H), 7.90 (d, J=8.4Hz, 1H), 7.65 (t, J=7.4Hz, 1H), 7.54 (t, J=7.5Hz, 1H), 7.40-7.33 (m, 5H), 5.39-5.33 (m 1H), 5.26-5.15 (m, 1H), 4.59 (d, J=13.5Hz, 1H), 4.47 (d, J=13.5Hz, 1H), 3.63 (dd, J=14.8Hz, J=10.6Hz, 1H), 3.42 (dd, J=14.8Hz, J=2.5Hz, 1H), 2.89 (s, 3H), 2.79 (s, 3H), 2.10-1.94 (m, 1H), 1.81-1.64 (m, 1H), 0.97 (t, J=7.2Hz, 3H); MS: (M⁺+1) 502.

- 10 (f) Morpholine-4-carboxylic acid (R)-1-[(S)-1-(1-benzylcarbamoyl-methanoyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester, (Compound 25).



- By proceeding in a manner similar to Example 4(a) above but using (R)-3-amino-2-hydroxy-pentanoic acid benzylamide TFA salt (Reference Example 19) in step 3 there was prepared morpholine-4-carboxylic acid (R)-1-[(S)-1-(1-benzylcarbamoyl-methanoyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester. ¹H NMR: (DMSO) 9.27 (t, J=5.5Hz, 1H), 8.67 (d, J=8.1Hz, 1H), 7.40-7.20 (m, 10H), 5.42-5.34 (m 1H), 4.96-4.85 (m, 1H), 4.64-4.24 (m, 4H), 3.66-3.28 (m, 10H), 1.90-1.72 (m, 1H), 1.63-1.46 (m, 1H), 0.89 (t, J=7.2Hz, 3H); MS: (M⁺+1) 560.

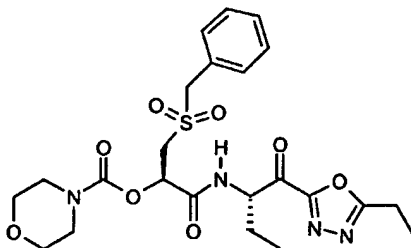
- 20 (g) Morpholine-4-carboxylic acid (S)-1-[(S)-1-(oxazol[4,5-b]pyridine-2-carbonyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester



-93-

By proceeding in a manner similar to Example 4(a) above but using (S)-2-amino-1-oxazolo[4,5-b]pyridin-2-yl-butan-1-ol TFA salt (Reference Example 20) there was prepared morpholine-4-carboxylic acid (S)-1-[(S)-1-(oxazolo[4,5-b]pyridine-2-carbonyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester. ¹H NMR: (DMSO) 9.00 (d, J=6.4Hz, 1H), 8.73 (m, 1H), 8.39 (d, J=8.4Hz, 1H), 7.69-7.64 (m, 1H), 7.45-7.30 (m, 5H), 5.37 (d, J=10.4Hz, 1H), 5.19-5.13 (m, 1H), 4.57 (d, J=13.6Hz, 1H), 4.46 (d, J=13.6Hz, 1H), 3.67-3.23 (m, 10H), 2.10-1.98 (m, 1H), 1.80-1.69 (m, 1H), 0.99 (t, J=7.0Hz, 3H). MS: (M+H)⁺ 545.

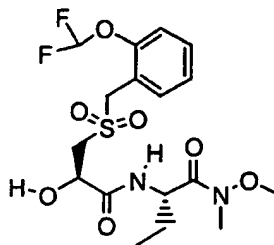
(h) Morpholine-4-carboxylic acid (S)-1-[(S)-1-(5-ethyl-[1,3,4]oxadiazole-2-carbonyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester



By proceeding in a manner similar to Example 4(a) above but using 2-amino-1-(5-ethyl-[1,3,4]oxadiazol-2-yl-butan-1-ol {Reference Example 11(m)} there was prepared morpholine-4-carboxylic acid (S)-1-[(S)-1-(5-ethyl-[1,3,4]oxadiazole-2-carbonyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester. ¹H NMR: (DMSO) 8.95 (d, J=6.0Hz, 1H), 7.41-7.33 (m, 5H), 5.35 (d, J=10.0Hz, 1H), 4.97-4.91 (m, 1H), 4.63-4.45 (m, 2H), 3.64-3.23 (m, 10H), 2.96 (q, J=7.2Hz, 2H), 1.99-1.89 (m, 1H), 1.75-1.64 (m, 1H), 1.30 (t, J=7.6Hz, 3H), 0.94 (t, J=7.2Hz, 3H). MS: (M+H)⁺ 523.

EXAMPLE 5

(S)-2-[(R)-3-[2-(1,1-Difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propanoylamino]-N-methoxy-N-methyl-butylamide, (Compound 32)



To a solution of (R)-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionic acid {1.24g, 4mmol, Reference Example 1(a)} in CH₂Cl₂ (20ml) was added HOBt (0.74g, 4.8mmol), EDC (1.15g, 6mmol), (R)-2-amino-N-methoxy-N-methyl-butylamide TFA salt (1.04g, 4mmol), prepared as in reference 2, and NMM (1.6g, 16mmol). After stirring for 14 hours at room temperature, the

-94-

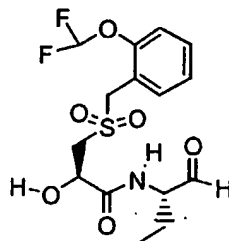
reaction mixture was diluted with 150ml of ethyl acetate. The mixture was washed with saturated NaHCO₃ and brine before drying with anhydrous MgSO₄. This crude product was then filtered, concentrated and purified by flash column chromatography using silica gel with hexane/ acetate as eluent to yield (S)-2-{(R)-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-

5 propanoylamino}-N-methoxy-N-methyl-butylamide (1.45g); ¹HNMR (CDCl₃): 7.6-7.5(d, J=7.67Hz, 1H), 7.5-7.35(m, 2H), 7.31-7.15(m, 2H), 6.63(t, J=73.4Hz, 1H), 5.0-4.85(br., 1H), 4.7-4.6(m, 1H), 4.55-4.48(m, 2H), 4.45-4.35(m, 1H), 3.80(s, 3H), 3.6-3.8(m, 1H), 3.35-3.2(m, 1H), 1.78(s, 3H), 2.0-1.5(m, 2H), 0.93(t, J=6.9Hz, 3H); MS: 437.44(M-1), 439.4(M+1).

10

EXAMPLE 6

(R)-3-[2-(1,1-Difluoro-methoxy)-phenylmethanesulfonyl]-N-((S)-1-formyl-propyl)-2-hydroxy-
propionamide. (Compound 23)



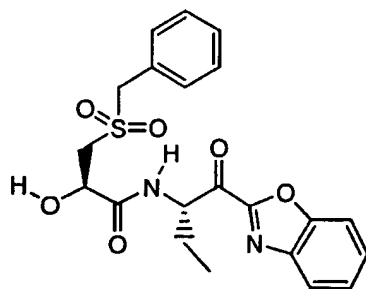
To a solution of (S)-2-{(R)-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-
15 propanoylamino}-N-methoxy-N-methyl-butylamide (1.3g, 3mmol, Example 5) in ethyl ether (50mL) at 0°C under N₂, was added 1N LAH solution of ethyl ether (3ml). After stirring for 3 hours at 0°C, 1ml of ethyl acetate and saturated NH₄Cl solution was added. The crude product was then extracted with ether, washed with brine, dried with MgSO₄, filtered and concentrated. The residue was purified by flash column chromatography using silica gel with hexane/ acetate as eluent to yield (R)-3-[2-(1,1-
20 difluoro-methoxy)-phenylmethanesulfonyl]-N-((S)-1-formyl-propyl)-2-hydroxy-propionamide (0.66g); ¹HNMR (DMSO): 9.43(s, 1H), 8.42(d, J=7.45Hz, 1H), 7.6-7.0(m, 4H), 7.12(t, J=73.93Hz, 1H), 6.52(d, J=6.45Hz, 1H), 5.2-5.17(m, 1H), 4.65-4.53(m, 2H), 4.12-4.0(m, 1H), 3.63-3.55(m, 2H), 1.7-1.4(m, 2H), 0.89(t, J=6.8Hz, 3H); MS: 378.2(M-1), 380.4(M+1).

25

EXAMPLE 7

(R)-N-[(S)-1-(1-benzooxazol-2-yl-methanoyl)-propyl]-2-hydroxy-3-phenyl-methanesulfonyl-
propionamide, (Compound 5)

-95-



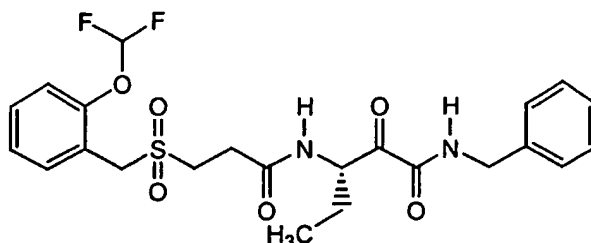
Step 1. To a solution of (R)-3-Phenylmethanesulfonyl-2-triisopropylsilanyloxy-propionic acid {556mg, 1mmol, Reference Example 3} in CH_2Cl_2 (10mL) at room temperature was added HOBT (183mg, 1.2mmol), EDC (288mg, 15mmol), (S)-2-Amino-1-benzooxazol-2-yl-butanol (206mg, 1mmol) and NMM (202mg, 2mmol). The mixture was then stirred overnight at room temperature before being diluted with ethyl acetate (100mL), washed with saturated NaHCO_3 , brine, dried with anhydrous MgSO_4 , filtered and concentrated. The crude product was then purified by flash column chromatography using silica gel with hexane/acetate as eluent (to yield 180mgs of product). This compound was dissolved in CH_2Cl_2 , Dess-Martin Periodinane (196mg, 0.46mmol) was added at room temperature and the mixture was stirred for 2 hours. Saturated $\text{Na}_2\text{S}_2\text{O}_3$ - NaHCO_3 solution (5mL) was added and stirred for a further 30 minute before extraction with ethyl acetate and washing sequentially with saturated NaHCO_3 solution and brine. The crude product was then dried with anhydrous MgSO_4 , filtered, concentrated and purified by flash column chromatography using silica gel with hexane/acetate as eluent to yield (R)-N-[(S)-1-(1-benzooxazol-2-yl-methanoyl)-propyl]-3-phenylmethanesulfonyl-2-triisopropylsilanyloxy-propionamide.

Step 2. (R)-N-[(S)-1-(1-Benzooxazol-2-yl-methanoyl)-propyl]-3-phenylmethanesulfonyl-2-triisopropylsilanyloxy-propionamide (120mg, 0.2mmol), in CH_3CN (10mL), 48% HF/ water solution (1mL) were mixed and stirred at room temperature for 16 hours. Saturated NaHCO_3 solution was added carefully to adjust the pH to between 8 and 9. The product was extracted with ethyl acetate (100mL), washed with brine and dried with magnesium sulfate. The solvent was removed and the product crystallized from acetate and hexane to yield (R)-N-[(S)-1-(1-benzooxazol-2-yl-methanoyl)-propyl]-2-hydroxy-3-phenyl-methanesulfonyl-propionamide as a white solid (85% yield); ^1H NMR: (DMSO) 8.29 (d, $J=7.9\text{Hz}$, 1H), 7.74 (d, $J=7.9\text{Hz}$, 1H), 7.59 (t, $J=8.1\text{Hz}$, 1H), 7.46-7.35 (m, 7H), 6.52 (d, $J=6.6\text{Hz}$, 1H), 5.08-4.99 (m, 1H), 4.58-4.47 (m, 3H), 3.35-3.28 (m, 2H), 2.05-1.90 (m, 1H), 1.81-1.65 (m, 1H), 0.91 (t, $J=7.2\text{Hz}$, 3H); MS: ($\text{M}^+ + 1$) 431.

EXAMPLE 8

(a) (S)-3-{3-[2-(1,1-Difluoro-methoxy)-phenylmethanesulfonyl]-propanoylamino}-2-oxo-pentanoic acid benzylamide, (Compound 27)

-96-



Step 1. A mixture of (R)-3-amino-2-hydroxy-pentanoic acid benzylamide TFA salt (70mg, 0.22mmol), 3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-propionic acid (64mg, 0.22mmol, Reference Example 19) HOBT (33mg, 0.22mmol), EDC (63mg, 0.325mmol), 1mL dichloromethane and N-methyl morpholine (48μL, 0.434mmol). The mixture was allowed to stir 16 hours. The product was extracted into 60mL ethyl acetate and washed with two 10mL portions of 1N HCl, 10mL water, and two 10mL portions of saturated NaHCO₃, dried over MgSO₄ and concentrated to give 105mg of crude (R)-3-{3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-propanoylamino}-2-hydroxy-pentanoic acid benzylamide (0.21mmol, 100% yield).

10

Step 2. To a 1mL dichloromethane solution of 105 mg of (R)-3-{3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-propanoylamino}-2-hydroxy-pentanoic acid benzylamide (0.21 mmol) was added Dess Martin periodinane (179mg, 0.42 mmol). The mixture was allowed to stir for 16 hours, then 10mL of 0.26M Na₂S₂O₃ in saturated NaHCO₃ was added and the mixture was extracted with two 30mL portions of ethyl acetate and washed with three 15mL portions of saturated NaHCO₃. The organic layer was dried over MgSO₄ and concentrated. The product was purified by silica gel chromatography using 3:1 hexane:ethyl acetate eluent and crystallized from diethyl ether and hexane to give (S)-3-{3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-propanoylamino}-2-oxo-pentanoic acid benzylamide (28mg, 0.054mmol, 26% yield); ¹H NMR: (CDCl₃) 7.0-7.47 (m, 9H), 6.49 (m, 1H), 6.24 (m, 1H), 5.22 (m, 1H), 4.40 (m, 2H), 4.30 (m, 3H), 3.23 (m, 2H), 2.70 (m, 2H), 2.01 (m, 1H), 1.68 (m, 1H), 0.85 (m, 3H); MS: (M⁺+1) 499.4, 496.53.

20

The following compounds were prepared by the method of Example 8:

N-[(S)-1-(1-Benzoxazol-2-yl-methanoyl)-propyl]-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-propionamide (Compound 26); ¹H NMR: (CDCl₃) 7.85 (d, J=7.6Hz, 1H), 7.7-7.0 (m, 7H), 6.51 (m, 2H), 5.60 (m, 1H), 4.34 (m, 3H), 3.29 (m, 2H), 2.80 (m, 2H), 2.13 (m, 1H), 1.87 (m, 1H), 0.96 (m, 3H); MS: (M⁺+1) 481, 480.48;

25

N-[(S)-1-(1-Benzoxazol-2-yl-methanoyl)-3-phenyl-propyl]-3-p-tolylmethanesulfonyl-propionamide (Compound 30); ¹H NMR: (CDCl₃) 7.9 (m, 1H), 7.62 (m, 1H), 7.56 (td, J=6.9, 1.2Hz, 1H), 7.47 (td, J=7.1, 1.2Hz, 1H), 7.3-7.1 (m, 9H), 6.47 (d, J=7.7Hz, 1H), 5.71 (m, 1H), 4.22 (s, 2H), 3.20 (m, 2H), 2.71 (m, 4H), 2.47 (m, 1H), 2.33 (s, 3H), 2.21 (m, 1H); MS: (M⁺+1) 505.2, 504.60.

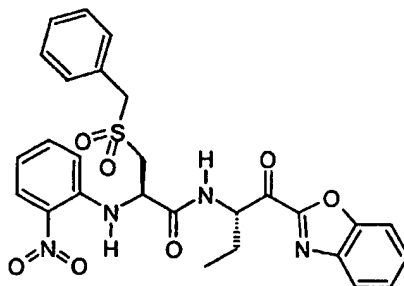
30

3-(2-Difluoromethoxy-phenylmethanesulfonyl)-N-(1-ethyl-2,3-dioxo-3-pyrrolidin-1-yl-propyl)-

-97-

propionamide;3-(2-Difluoromethoxy-phenylmethanesulfonyl)-N-(1-ethyl-3-morpholin-4-yl-2,3-dioxo-propyl)-propionamide;3-(2-Difluoromethoxy-phenylmethanesulfonyl)-N-(1-ethyl-2,3-dioxo-3-piperazin-1-yl-propyl)-5 propionamide;3-(2-Difluoromethoxy-phenylmethanesulfonyl)-N-[3-(1,1-dioxo-1,6-thiomorpholin-4-yl)-1-ethyl-2,3-
dioxo-propyl]-propionamide;3-(2-Difluoromethoxy-phenylmethanesulfonyl)-N-[1-ethyl-3-(4-methyl-sulfonyl-piperazin-1-yl)-2,3-
dioxo-propyl]-propionamide;10 3-[3-(2-Difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid
dimethylamide;3-[3-(2-Difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid cyclopentyl-
ethyl-amide;3-[3-(2-Difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid15 phenylamide;3-[3-(2-Difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid pyridin-3-
ylamide;3-[3-(2-Difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid (tetrahydro-
pyran-4-yl)-amide;20 3-[3-(2-Difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid (1-benzoyl-
piperidin-4-yl)-amide; and3-[3-(2-Difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid (2-
morpholin-4-yl-ethyl)-amide.

25

EXAMPLE 9(R)-N-[(S)-1-(1-Benzooxazol-2-yl-methanoyl)-propyl]-2-(2-nitro-phenylamino)-3-
phenylmethanesulfonyl-propionamide, (Compound 28)

Step 1. 3-Benzylsulfanyl-2-(2-nitro-phenylamino)-propionic acid (350mg, 1.05 mmol, Reference
 30 Example 5) was dissolved in 20mL methanol, treated with a 20mL aqueous solution of Oxone®

-98-

(970mg, 0.12mmol), and stirred for 72 hours. Water (300mL) was added and the precipitate was filtered and dried to give 2-(2-nitro-phenylamino)-3-phenylmethanesulfonyl-propionic acid (215mg, 0.59mmol, 56%yield)

5 Step 2. A mixture of 2-(2-nitro-phenylamino)-3-phenylmethanesulfonyl-propionic acid (215mg, 0.59mmol), HOBT (136mg, 0.148mmol), EDC (408mg, 2.13mmol), (S)-2-amino-1-benzooxazol-2-yl-butan-1-ol (122mg, 0.59mmol, {Reference Example 17(a)}), 2.4mL dichloromethane and N-methyl morpholine (97 μ L, 0.89mmol) was allowed to stir 16 hours. The product was extracted into 20mL ethyl acetate and washed with three 5mL portions of 1N HCl, and one 30mL portion of saturated
10 NaHCO₃, dried over MgSO₄ and concentrated to give (R)-N-[(S)-1-(1-benzooxazol-2-yl-1-hydroxy-methyl)-propyl]-2-(2-nitro-phenylamino)-3-phenylmethane-sulfonyl-propionamide (223mg, 0.40mmol, 45% yield).

Step 3. (R)-N-[(S)-1-(1-Benzooxazol-2-yl-1-hydroxy-methyl)-propyl]-2-(2-nitro-phenylamino)-3-
15 phenylmethane-sulfonyl-propionamide (223mg, 0.4mmol) was dissolved in 1.6mL dichloromethane and treated with Dess Martin periodinane (342mg, 0.80 mmol). The mixture was allowed to stir for 16 hours, then 20mL of 0.26M Na₂S₂O₃ in saturated NaHCO₃ was added and the mixture was extracted with two 30mL portions of ethyl acetate and washed with three 5mL portions of saturated NaHCO₃. The organic layer was dried over MgSO₄ and concentrated. The crude product was dissolved in a
20 minimum amount of hot ethyl acetate and crystallized by addition of dry diethyl ether. This crystallization was repeated to give clean (R)-N-[(S)-1-(1-Benzooxazol-2-yl-methanoyl)-propyl]-2-(2-nitro-phenylamino)-3-phenylmethanesulfonyl-propionamide (97mg, 0.176mmol, 44% yield); ¹H NMR: (DMSO) 8.67 (m, 1H), 8.12 (m, 1H), 7.81 (m, 1H), 7.65-7.35 (m, 10H), 6.78 (m, 2H), 5.51 (m, 1H), 4.68 (m, 1H), 4.37 (s, 2H), 3.62 (m, 1H), 3.38 (m, 1H), 2.15 (m, 1H), 1.91 (m, 1H), 0.98 (m, 3H);
25 MS: (M⁺+1) 551.0, 550.58.

The following compound was prepared by the method of Example 9:

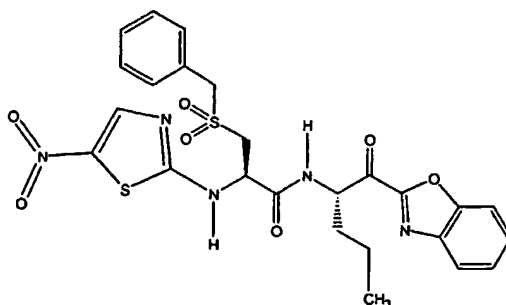
N-[1-(Benzooxazole-2-carbonyl)-propyl]-3-phenylmethanesulfonyl-2-(pyrimidin-2-ylamino)-propionamide.

30

EXAMPLE 10

(R)-N-[(S)-1-(1-Benzooxazol-2-yl-methanoyl)-butyl]-2-(5-nitro-thiazol-2-ylamino)-3-phenylmethanesulfonyl-propionamide, (Compound 29)

-99-



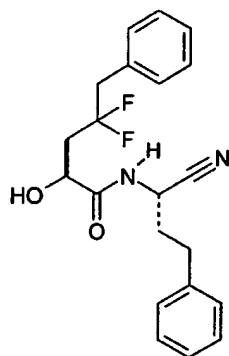
Step 1. A mixture of (R)-3-benzylsulfanyl-2-(5-nitro-thiazol-2-ylamino)-propionic acid (42mgmg, 0.123mmol, Reference Example 6) HOBT (28mg, 0.148mmol), EDC (29mg, 0.148mmol), (S)-2-amino-1-benzooxazol-2-yl-pentan-1-ol {27mg, 0.123mmol, Reference Example 17(c)}, 1mL
 5 dichloromethane and N-methyl morpholine (14μL, 0.123mmol) was allowed to stir for 16 hours. The product was extracted into 60mL ethyl acetate and washed with one 30mL portion of 1N HCl, and one 30mL portion of saturated NaHCO₃, dried over MgSO₄ and concentrated to give (R)-N-[(S)-1-(1-benzooxazol-2-yl-1-hydroxy-methyl)-butyl]-3-benzylsulfanyl-2-(5-nitro-thiazol-2-ylamino)-propionamide (41.8mg, 0.077mmol, 63% yield).

Step 2. (R)-N-[(S)-1-(1-Benzooxazol-2-yl-1-hydroxy-methyl)-butyl]-3-benzylsulfanyl-2-(5-nitro-thiazol-2-ylamino)-propionamide (41.8mg, 0.077mmol) was dissolved in 0.5mL methanol, treated with a 0.5mL aqueous solution of Oxone® (43mg, 0.069mmol), and stirred for 1 hour. Methanol was removed under reduced pressure and 2mL water was added. The mixture was extracted with two
 15 10mL portions of ethyl acetate, dried over MgSO₄, and concentrated. It was then dissolved in 0.5mL dichloromethane and treated with Dess Martin periodinane (65mg, 0.154 mmol). The mixture was allowed to stir for 16 hours, then 5mL of 0.26M Na₂S₂O₃ in saturated NaHCO₃ was added and the mixture was extracted with two 10mL portions of ethyl acetate and washed with three 5mL portions of saturated NaHCO₃. The organic layer was dried over MgSO₄ and concentrated. The product was
 20 purified by triturating with diethyl ether to give (R)-N-[(S)-1-(1-benzooxazol-2-yl-methanoyl)-butyl]-2-(5-nitro-thiazol-2-ylamino)-3-phenylmethanesulfonyl-propionamide (28mg, 0.054mmol, 26% yield);
¹H NMR: (CDCl₃) 7.96 (s, 1H), 7.87 (m, 1H), 7.7-7.3 (m, 9H), 5.57 (m, 1H), 5.06 (m, 1H), 4.47 (m, 2H), 3.75 (m, 1H), 3.48 (m, 1H), 2.09 (m, 1H), 1.85 (m, 1H), 1.43 (m, 1H), 1.24 (m, 1H), 0.94 (m, 3H); MS: (M⁺+1) 572.2, 571.63.

EXAMPLE 11

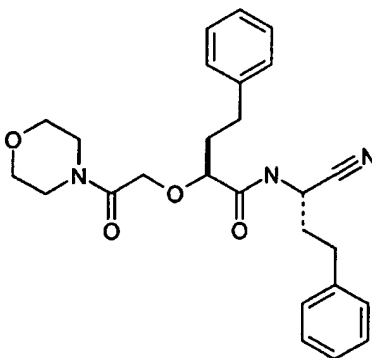
(a) (2S) (4,4-Difluoro-2-hydroxy-5-phenyl-pentanoic acid (1(S)-cyano-3-phenyl-propyl)-amide,
 (Compound 33)

-100-



To a mixture of amino-acetonitrile hydrochloride (0.37 mmol, 72.6mg), (2S)-4,4-difluoro-2-hydroxy-5-phenyl-pentanoic acid (1.0 eq., 0.37 mmol, 85.0mg, Reference Example 7) and N,N-diisopropylethylamine (2.2 eq., 0.81 mmol, 105.2mg) in dry dichloromethane (4 mL) under nitrogen was added PyBOP® (1.1 eq., 0.41 mmol, 212mg). The mixture was stirred at room temperature for 15.5 hours and then concentrated in vacuum. The residue was diluted with ethyl acetate (30ml) and the mixture was washed with water (30mL), then with sodium bicarbonate (30mL) and then with water (30mL). The organic layer was dried over MgSO₄ and then concentrated in vacuum. The residue was purified over 10g silica gel, eluting with a mixture of ethyl acetate and heptane (1:2, v/v) to afford (2S)-4,4-difluoro-2-hydroxy-5-phenyl-pentanoic acid (1S)-cyano-3-phenyl-propyl-amide as a light tan solid (67.4 mg, 48.9%). ¹H NMR (CDCl₃) 7.3 (m, 10H), 7.1 (d, J=7 Hz, 1H), 4.8 (q, J=7.4 Hz, 1H), 4.53 (bd, J=9.5 Hz, 1H), 3.2 (dt, J=16.2, 4.2 Hz, 2H), 2.96 (s, 1H), 2.85 (m, 2H), 2.5 (m, 1H), 2.3-0.9 (m, 3H). LC/MS 89% parent (M+1).

(b) N-(1S)-cyano-3-phenyl-propyl)-2-(S)-(2-morpholin-4-yl-2-oxo-ethoxy)-4-phenyl-butyramide, (Compound 34)

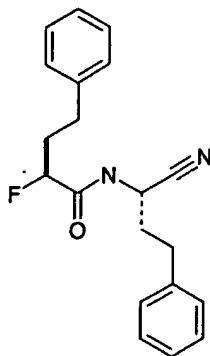


By proceeding in a manner similar to Example 11(a) above but using (S)-2-amino-4-phenyl-butyronitrile hydrochloride and 2-(S)-(2-morpholin-4-yl-2-oxo-ethoxy)-4-phenyl-butyric acid [Reference Example 8] there was prepared N-(1S)-cyano-3-phenyl-propyl)-2-(S)-(2-morpholin-4-yl-2-oxo-ethoxy)-4-phenyl-butyramide as an oil. ¹H NMR (CDCl₃) 9.4 (d, J=8.2 Hz, 1H), 7.3 (m, 10H),

-101-

4.75 (q, $J=7.5$ Hz, 1H), 4.63 (d, $J=15.1$ Hz, 1H), 3.95 (d, $J=15.3$ Hz, 1H), 3.87 (dd, $J=8.2, 3.9$ Hz, 1H), 3.7 (m, 6H), 3.32 (m, 2H), 2.85 (m, 4H), 2.1 (m, 3H), 2.05 (m, 1H). LC/MS 100% ($M+1$) 450.

(c) N-(1-(S)-cyano-3-phenyl-propyl)-2-(S)-fluoro-4-phenyl-butyramide, (Compound 35)

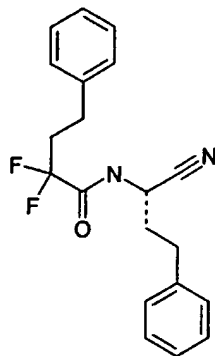


5

By proceeding in a manner similar to Example 11(a) above but using (S)-2-amino-4-phenyl-butyronitrile hydrochloride and (2S)-2-fluoro-4-phenyl-butyric acid (Reference Example 9) there was prepared N-(1-(S)-cyano-3-phenyl-propyl)-2-(S)-fluoro-4-phenyl-butyramide as a light tan solid. ^1H NMR (CDCl_3) 7.3 (m, 10H), 6.6 (bs, 1H), 4.95 (ddd, $J=49.2, 8.2, 3.5$ Hz, 1H), 4.8 (m, 1H), 3.8 (m, 4H), 2.3 (m, 1H), 2.2 (m, 3H). MS (CI, $M+1$) 325.

10

(d) N-(1-(S)-cyano-3-phenyl-propyl)-2,2-difluoro-4-phenyl-butyramide, (Compound 36)



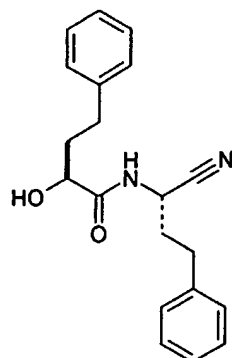
By proceeding in a manner similar to Example 11(a) above but using (S)-2-amino-4-phenyl-butyronitrile hydrochloride and 2,2-difluoro-4-phenyl-butyric acid there was prepared N-(1-(S)-cyano-3-phenyl-propyl)-2,2-difluoro-4-phenyl-butyramide as a white solid. ^1H NMR (CDCl_3) 7.3 (m, 10H), 6.6 (d, $J=8.1$ Hz, 1H), 4.83 (q, $J=7.4$ Hz, 1H), 2.88 (dt, $J=7.5, 2.5$ Hz, 2H), 2.79 (t, $J=8$ Hz, 2H), 2.4 (m, 2H), 2.2 (q, $J=7.5$ Hz, 2H). LC/MS 50% ($M+1$) 343.

15

(e) N-(1-(S)-cyano-3-phenyl-propyl)-2-(S)-hydroxy-4-phenyl-butyramide, (Compound 37)

20

-102-

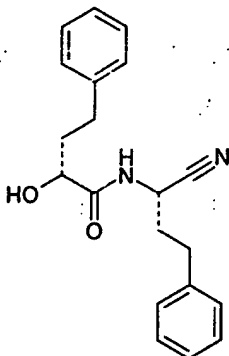


By proceeding in a manner similar to Example 11(a) above but using (S)-2-amino-4-phenyl-butyronitrile hydrochloride and (2S)-2-hydroxy-4-phenyl-butyric acid there was prepared

N-(1-(S)-cyano-3-phenyl-propyl)-2-(S)-hydroxy-4-phenyl-butyramide as a white solid. ¹H NMR

(CDCl₃) 7.3 (m, 10H), 6.9 (d, J=8.4 Hz, 1H), 4.86 (q, J=7.4 Hz, 1H), 4.2 (m, 1H), 2.84 (t, J=7.1 Hz, 2H), 2.77 (t, J=7.8 Hz, 2H), 2.5 (d, J=4.7 Hz, H), 2.2 (m, 3H), 1.95 (m, 1H). LC/MS 49% (M+1) 323.

(f) N-(1-(S)-cyano-3-phenyl-propyl)-2-(R)-hydroxy-4-phenyl-butyramide, (Compound 38)

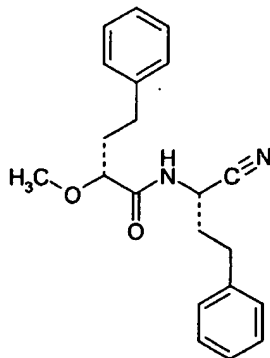


By proceeding in a manner similar to Example 11(a) above but using (S)-2-amino-4-phenyl-

butyronitrile hydrochloride and (2R)-2-hydroxy-4-phenyl-butyric acid there was prepared N-(1-(S)-cyano-3-phenyl-propyl)-2-(R)-hydroxy-4-phenyl-butyramide as a white solid. ¹H NMR (CDCl₃) 7.4-7.1 (m, 10H), 6.9 (d, J=8.7 Hz, 1H), 4.87 (q, J=7.3 Hz, 1H), 4.1 (m, 1H), 2.85 (t, J=7.5 Hz, 2H), 2.77 (t, J=8.4 Hz, 2H), 2.3 (d, J=5.1 Hz, 1H), 2.2 (m, 3H), 2.0 (m, 1H). LC/MS 94% (M+1) 323.

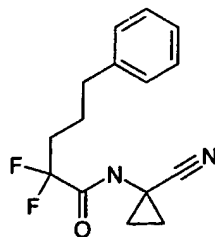
-103-

- (g) N-(1-(S)-cyano-3-phenyl-propyl)-2-(R)-methoxy-4-phenyl-butyramide, (Compound 39)



- By proceeding in a manner similar to Example 11(a) above but using (S)-2-amino-4-phenyl-butyronitrile hydrochloride (0.407 mmol, 80mg) and 2(R)-methoxy-4-phenyl-butyric acid (Reference Example 10) there was prepared N-(1-(S)-cyano-3-phenyl-propyl)-2-(R)-methoxy-4-phenyl-butyramide as a white solid (91.8mg, 67%). ¹H NMR (CDCl₃) 7.2 (m, 10H), 6.8 (d, J=8.5 Hz, 1H), 4.86 (q, J=7.5 Hz, 1H), 3.67 (dd, J=6.5, 4.5 Hz, 1H), 3.35 (s, 3H), 2.85 (m, 2H), 2.68 (t, J=8.0 Hz, 2H), 2.2-2.0 (m, 4H). LC/MS 84% (M⁺1) 337.

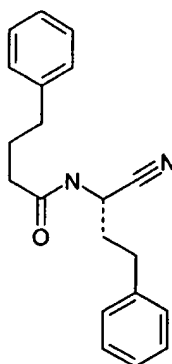
- 10 (h) 2,2-Difluoro-5-phenyl-pentanoic acid (1-cyano-cyclopropyl)-amide, (Compound 40)



- By proceeding in a manner similar to Example 11(a) above but using 2,2-difluoro-5-phenyl-pentanoic acid and 1-amino-cyclopropanecarbonitrile hydrochloride there was prepared 2,2-difluoro-5-phenyl-pentanoic acid (1-cyano-cyclopropyl)-amide. ¹H NMR (CDCl₃) δ 1.32 (m, 2H), 1.64 (m, 2H), 1.82 (m, 2H), 2.12 (m, 2H), 2.8-2.56 (m, 2H), 6.82 (m, 1H), 7.36-7.15 (m, 5H). MS (ES-) 277.

- (i) N-(1-(S)-Cyano-3-phenyl-propyl)-4-phenyl-butyramide, (Compound 41)

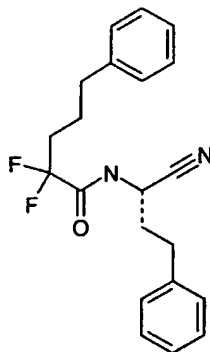
-104-



By proceeding in a manner similar to Example 11(a) above but using (S)-2-amino-4-phenyl-butyronitrile hydrochloride and 4-phenylbutyric acid there was prepared N-(1-(S)-cyano-3-phenyl-propyl)-4-phenyl-butyramide as a colorless oil. ¹H NMR (CDCl₃): δ 7.3 (m, 10H), 6.0 (d, J=8.3 Hz, 1H), 4.9 (q, J=7.4 Hz, 1H), 2.8 (m, 2H), 2.65 (t, J=7.4 Hz, 2H), 2.15 (m, 4H), 1.95 (m, 2H). LC/MS 100% (M+1) 307.

EXAMPLE 12

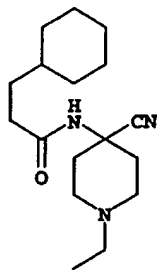
10 2,2-difluoro-5-phenyl-pentanoic acid ((S)-1-cyano-3-phenyl-propyl)-amide, (Compound 42)



A mixture of 2,2-difluoro-5-phenyl-pentanoic acid (109mg, 0.509 mmol), (S)-2-amino-4-phenyl-butyronitrile hydrochloride (103mg, 0.523 mmol) and HATU (206mg, 0.542 mmol) in DMF (4mL) and stirred at room temperature for 5 hours then evaporated under reduced pressure. The residue was taken in ethyl acetate washed with 1N HCl, sodium bicarbonate and then water. Organic extract was dried over Na₂SO₄ and then evaporated under vacuum to give orange oil. The residue was subjected to mpls, eluting with a mixture of ethyl acetate and heptane (1:9, v/v) to give 2,2-difluoro-5-phenyl-pentanoic acid ((S)-1-cyano-3-phenyl-propyl)-amide as colorless oil (82 mg). ¹H NMR (CDCl₃) 7.3-7.1 (m, 10H), 6.9 (bs, 1H), 4.80 (q, J=7.5 Hz, 1H), 2.80 (dt, J=7.3, 2.7 Hz, 2H), 2.65 (t, J=7.5 Hz, 2H), 2.2-2.0 (m, 4H), 1.8 (m, 2H). MS 357 (MH⁺), 379 (M+Na).

EXAMPLE 13

-105-

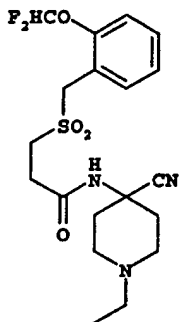
(a) N-(4-Cyano-1-ethyl-piperidin-4-yl)-3-cyclohexyl-propionamide

Step 1. To a stirred solution of 1-ethyl-4-piperidone (25g, 0.197mol) in 300ml of diethyl ether, and NH_4Cl (22.3g, 0.41mol), was added NaCN (14.5g, 0.295mol, in 70ml water) drop-wise at room temperature. After stirring for 24h the diethyl ether was separated and the water phase was extracted with n-BuOH, then washed with brine and dried. After removal of most of the n-BuOH under reduced pressure, the residue was diluted with 50ml of diethyl ether and then acidified with 2N HCl in diethyl ether solution at 0°C. The solid was dried under vacuum to yield 45g of 4-amino-1-ethyl-piperidine-4-carbonitrile HCl salt.

10

Step 2. To a stirred mixture of 3-cyclohexyl-propionic acid (156mg, 1mmol), 4-amino-1-ethyl-piperidine-4-carbonitrile HCl salt (227, 1mmol, prepared as in step 1 above), and HATU (570mg, 1.5mmol) in MeCl_2 (5ml), was added N,N-diisopropylethylamine (516mg, 4mmol) at room temperature. After stirring for 14 hours, the reaction mixture was extracted with ethyl acetate. The organic layer was washed with saturated NaHCO_3 , brine, dried with MgSO_4 and concentrated to yield N-(4-Cyano-1-ethyl-piperidin-4-yl)-3-cyclohexyl-propionamide (170mg). LC-MS: elution time = 2.25min. 290.2(M-1), 292.2(M+1). (MS: API 150EX. LC: HP Agilent 1100 Series. Column: Phenomenex, 5u ODS3 100A 100X3mm.; Flow Rate: 2ml/min. Two solvent gradient: Solvent A, 99% water, 1% acetonitrile, 0.1% AcOH. Solvent B, 99% acetonitrile, 1% water, 0.1% AcOH. Gradient from 100% A, 0% B to 0% A, 100% B from t = 0 to t = 6min. Then gradient back to 100% A, 0% B from t = 7 to t = 15 min.).

20

(b) N-(4-Cyano-1-ethyl-piperidin-4-yl)-3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionamide

25

-106-

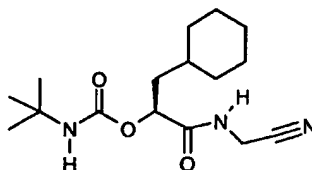
By proceeding in a similar manner to Example 13(a) but using 3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionic acid (294mg, 1mmol) and 4-amino-1-ethyl-piperidine-4-carbonitrile HCl salt(227, 1mmol) there was N-(4-cyano-1-ethyl-piperidin-4-yl)-3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionamide 260mg). LC-MS: R_T = 1.96min., 428.2(M-1), 430.3(M+1).

5 MS: API 150EX. (LC: Agilent 1100Series, Column: Phenomenex, 5u ODS3 100A 100X3mm. Flow Rate: 2ml/min. Two solvent gradient: Solvent A, 99% water, 1% acetonitrile, 0.1% AcOH. Solvent B, 99% acetonitrile, 1% water, 0.1% AcOH. Gradient from 100% A, 0% B to 0% A, 100% B from t = 0 to t = 6min. Then gradient back to 100% A, 0% B from t = 7 to t = 15 min.).

10

EXAMPLE 14

(S)-tert-Butyl-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester

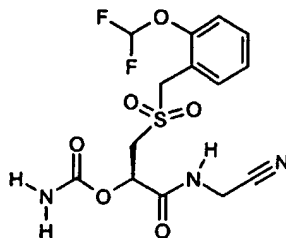


(S)-N-Cyanomethyl-3-cyclohexyl-2-hydroxy-propionamide (53mg, 0.252mmol) was dissolved in dichloromethane (1mL). Triethylamine (0.1mL) was added and then tert.-butyl isocyanate (0.034mL, 0.3mmol). The mixture was stirred at room temperature overnight. After dilution with ethyl acetate (100mL), the solution was washed with 1N aqueous. HCl, brine, sat. aqueous NaHCO₃, and brine, dried with MgSO₄ and evaporated under vacuum. Flash chromatography on silica gel (hexane/ethyl acetate 1:1) gave (S)-tert-Butyl-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester (63mg, 0.204mmol) as a white solid.

20

EXAMPLE 15

(a) (R)-Carbamic acid 1-(cyanomethyl-carbamoyl)-2-(2-difluoromethoxy-phenylmethanesulfonyl)-ethyl ester



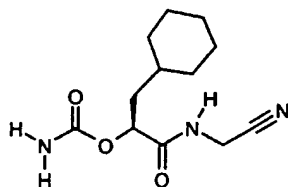
25 (R)-N-Cyanomethyl-3-(2-(1,1-difluoromethoxy)-phenylmethanesulfonyl)-2-hydroxy-propionamide {100mg, 0.287mmol, Example 1(a)} was dissolved in dichloromethane (2mL) and THF (1mL). Trichloroacetyl isocyanate (0.051mL, 0.43mmol) was added and the mixture was stirred for 1h. The

-107-

solvents were removed under vacuum and the residue was dissolved in 1,4-dioxane (10mL). 1N aqueous. HCl (5mL) was added and the mixture was heated at 70°C for 4h. After cooling to room temperature, the mixture was extracted with ethyl acetate. The combined organic layers were washed with brine, dried with MgSO₄ and evaporated under vacuum. Flash chromatography on silica gel (hexane/ ethyl acetate 1:3) gave (R)-carbamic acid 1-(cyanomethyl-carbamoyl)-2-(2-difluoromethoxy-phenylmethanesulfonyl)-ethyl ester (35mg, 0.089mmol) as a white solid. ¹H NMR: (DMSO) 8.90 (t, J=4.8Hz, 1H), 7.48-7.43 (m, 2H), 7.30-7.21 (m, 2H), 7.11 (t, J_{H,F}=73.6Hz, 1H), 6.98-6.76 (br, 2H), 5.28-5.23 (m, 1H), 4.60 (s, 2H), 4.15 (d, J=4.8Hz, 2H), 3.70 (dd, J=10.0Hz, J=14.8Hz, 1H), 3.54 (d, J=14.4Hz, 1H). MS: (M+H)⁺ 392.

10

(b) (S)-Carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester



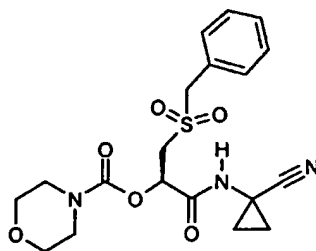
15

By proceeding in a manner similar to Example 8(a) above but using (R)-N-cyanomethyl-3-cyclohexyl-2-hydroxy-propionamide there was prepared (S)-Carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester. ¹H NMR: (DMSO) 8.63 (t, J=5.6Hz, 1H), 6.63 (br, 2H), 4.81-4.77 (m, 1H), 4.11 (d, J=5.2Hz, 2H), 1.74-0.81 (m, 13H). MS: (M+H)⁺ 254.

EXAMPLE 16

20

(a) (R)-Morpholine-4-carboxylic acid 1-(1-cyano-cyclopropylcarbamoyl)-2-phenylmethanesulfonyl-ethyl ester



25

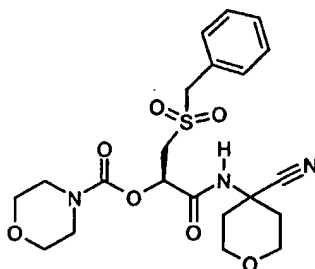
DMF was added to a mixture of (R)-morpholine-4-carboxylic acid 1-carboxy-2-phenylmethanesulfonyl-ethyl ester {from step 2 in Example 4(a)} (60mg, 0.168mmol), HATU (200mg, 0.52mmol), and 1-amino-cyclopropanecarbonitrile hydrochloride (100mg, 0.84mmol). 4-Methylmorpholine (0.5mL) was added and the mixture was stirred overnight. The mixture was diluted with ethyl acetate (100mL), washed with 1N aqueous. HCl, brine, sat. aqueous. NaHCO₃, brine, dried with MgSO₄ and evaporated under vacuum. Flash chromatography on silica gel (hexane/ethyl acetate

-108-

1:2) gave (R)-morpholine-4-carboxylic acid 1-(1-cyano-cyclopropylcarbamoyl)-2-phenylmethanesulfonyl-ethyl ester (7mg, 0.017mmol). ¹H NMR: (DMSO) 9.16 (s, 1H), 7.40-7.32 (m, 5H), 5.24-5.19 (m, 1H), 4.55 (d, J=13.2Hz, 1H), 4.48 (d, J=13.2Hz, 1H), 3.63-3.25 (m, 10H), 1.51-1.39 (m, 2H), 1.20-1.07 (m, 2H). MS: (M+H)⁺ 422.

5

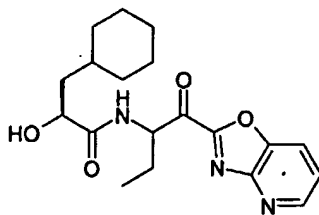
(b) (R)-Morpholine-4-carboxylic acid 1-(4-cyano-tetrahydro-pyran-4-ylcarbamoyl)-2-phenylmethanesulfonyl-ethyl ester



By proceeding in a manner similar to Example 16(a) above but using 4-amino-tetrahydropyran-4-carbonitrile hydrochloride {prepared according to Example 13(a) step1 but using tetrahydropyran-4-one} there was prepared (R)-morpholine-4-carboxylic acid 1-(4-cyano-tetrahydro-pyran-4-ylcarbamoyl)-2-phenylmethanesulfonyl-ethyl ester. LC-MS: elution time = 3.20min. 464.4(M-1), 466.2(M+1). (MS: API 150EX. LC: HP Agilent 1100 Series. Column: Phenomenex, 5u ODS3 100A 100X3mm.; Flow Rate: 2ml/min. Two solvent gradient: Solvent A, 99% water, 1% acetonitrile, 0.1% AcOH. Solvent B, 99% acetonitrile, 1% water, 0.1% AcOH. Gradient from 100% A, 0% B to 0% A, 100% B from t = 0 to t = 6min. Then gradient back to 100% A, 0% B from t = 7 to t = 15 min.)

EXAMPLE 17

3-Cyclohexyl-2-hydroxy-N-[1-(oxazolo[4,5-b]pyridine-2-carbonyl)-propyl]-propionamide



20

Step 1. To a stirred solution of [1-(hydroxy-oxazolo[4,5-b]pyridin-2-yl-methyl)-propyl]-carbamic acid *tert*-butyl ester (3.11g, 10mmol, prepared as described in Reference Example 20 step2.) in dioxane (4ml) was added HCl (4N solution in 5ml of dioxane) at room temperature. After 2 hours, ethyl ether(50ml) was added and the reaction mixture was filtered. The resultant solid was washed with an additional 20ml of ethyl ether and dried under vacuum to yield 3g of 2-amino-1-oxazolo[4,5-b]pyridin-2-yl-butan-1-ol HCl salt.

25

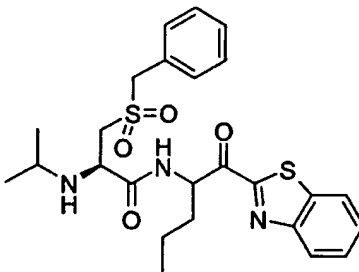
-109-

Step 2. To a stirred mixture of 3-cyclohexyl-2-hydroxy-propionic acid (155mg, 0.9mmol), 2-amino-1-oxazolo[4,5-*b*]pyridin-2-yl-butan-1-ol HCl salt, and HOBt (168mg, 1.1mmol) in MeCN (5ml), was added EDC (270mg, 1.4mmol) and N-methylmorpholine (0.45ml) at 23°C. After stirring for 14 hours, the reaction mixture was extracted with ethyl acetate. The organic layer was washed with saturated NaHCO₃, brine, dried with MgSO₄ and concentrated to yield 293 mg of 3-cyclohexyl-2-hydroxy-N-[1-(hydroxy-oxazolo[4,5-*b*]pyridin-2-yl-methyl)-propyl]-propionamide, which was used in step 3 following without further purification. MS: 360.3(M-1), 362.3(M+1), 384.2(M+Na).

Step 3. To a stirred solution of 3-cyclohexyl-2-hydroxy-N-[1-(hydroxy-oxazolo[4,5-*b*]pyridin-2-yl-methyl)-propyl]-propionamide (300mg, 0.83mmol) in MeCl₂(20ml), was added MnO₂(1.44g, 16.6mmol) at room temperature. After stirring for 30min. the mixture was filtered to remove MnO₂, and washed with 20ml of MeCl₂. The solvent was removed under vacuum and the residue was purified by silica gel column chromatography to yield 3-cyclohexyl-2-hydroxy-N-[1-(oxazolo[4,5-*b*]pyridine-2-carbonyl)-propyl]-propionamide (40mg). ¹H NMR (DMSO-*d*₆): 8.71(1H, dd, NH, diastereomer), 8.38(1H, dd,), 8.28(1H, m), 7.7-7.6(1H, m), 5.5-5.4(1H, m), 5.2-5.1(1H, m), 3.95-3.99(1H, br., OH), 2.1-1.95(1H, m), 1.85-1.75(1, m), 1.7-0.8(16H, m). MS: 358.1 (M-1), 360.1 (M+1), 382(M+Na).

EXAMPLE 18

(a) (R)-N-[1-(Benzothiazole-2-carbonyl)-butyl]-2-isopropylamino-3-phenylmethanesulfonyl-propionamide

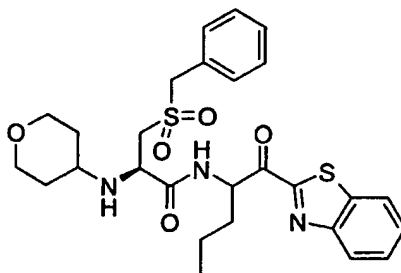


A solution of (R)-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-2-isopropylamino-3-phenylmethanesulfonyl-propionamide {30mg, 0.06mmol, Example 30(a)} in dichloromethane (10mL) was treated with Dess-Martin-periodinane (51mg, 0.12mmol). This mixture was stirred at room temperature for 45 minutes then treated with resin-bound thiosulfate (400mg, 0.6mmol) and stirring was continued for a further 24 hours then the mixture was treated with AP-Trisamine (270mg, 0.6mmol). After stirring for a further 24 hours the reaction mixture was filtered and the filtrate was evaporated to give (R)-N-[1-(benzothiazole-2-carbonyl)-butyl]-2-isopropylamino-3-phenylmethanesulfonyl-propionamide (23mg, 75%) as mixture of diastereomers. ¹H NMR (CDCl₃, 300MHz): 8.29-8.27 (m, 1H), 8.23-8.19 (m, 1H), 8.01-7.98 (m, 1H), 7.63-7.36 (m, 7H), 5.80-5.74 (m,

-110-

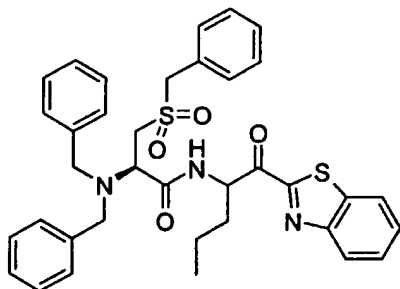
1H), 4.36-4.31 (m, 2H), [3.79 (dd, J=9.5Hz, 3Hz), 3.73 (dd, J=9Hz, 2.5Hz) 1H], 3.41-3.34 (m, 1H), 3.20-3.01 (m, 1H), 2.89-2.85 (m, 1H), 2.17-2.06 (m, 1H), 1.88-1.78 (m, 1H), 1.52-1.25 (m, 3H), 1.12-1.06 (m, 6H), [0.96 (t, J=7.5Hz) 0.95 (t, J=7.5Hz) 1H]. LC/MS m/z=502 (M+H).

- 5 (b) (R)-N-[1-(Benzothiazole-2-carbonyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide



- By proceeding in a similar manner to Example 18(a) but using (R)-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide {0.11 mmol, Example 29(b)} and subjecting the crude product to HPLC there was prepared (R)-N-[1-(benzothiazole-2-carbonyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide (10mg, 16%). LC/MS retention time 2.92min (TIC), m/z=544 (M+H) (determined with method A).

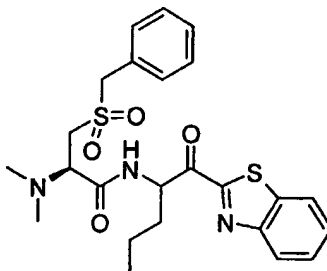
- 15 (c) (R)-N-[1-(Benzothiazole-2-carbonyl)-butyl]-2-dibenzylamino-3-phenylmethanesulfonyl-propionamide



- By proceeding in a similar manner to Example 18(a) but using (R)-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-2-dibenzylamino-3-phenylmethanesulfonyl-propionamide {0.11 mmol, Example 29(a)} and subjecting the crude product to HPLC there was prepared (R)-N-[1-(benzothiazole-2-carbonyl)-butyl]-2-dibenzylamino-3-phenylmethanesulfonyl-propionamide (4mg) as mixture of diastereomers. ¹H NMR (CDCl₃, 300MHz): 8.33-7.89 (m, 3H), 7.61-7.55 (m, 2H), 7.47-7.29 (m, 15H), 5.75 (m, 1H), [4.54 (d, J=14Hz), 4.51 (d, J=13.5Hz), 1H], [4.27 (d, J=14Hz), 4.25 (d, J=13.5Hz), 1H], 4.11-3.95 (m, 2H), [3.78 (d, J=13Hz), 3.76 (d, J=13Hz), 2H], [3.51 (d, J=13Hz), 3.50 (d, J=13Hz), 2H], 3.19-3.13 (m, 1H), 2.10-1.77 (m, 2H), 1.51-1.37 (m, 2H), 0.91-0.84 (m, 3H). LC/MS m/z=640 (M+H).

-111-

- (d) (R)-N-[1-(Benzothiazole-2-carbonyl)-butyl]-2-dimethylamino-3-phenylmethanesulfonyl-propionamide



5

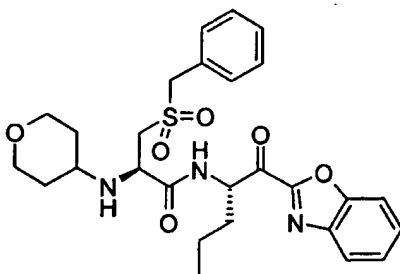
By proceeding in a similar manner to Example 18(a) but using (R)-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-2-dimethylamino-3-phenylmethanesulfonyl-propionamide {30mg, 0.06mmol, Example 30(b)}, and subjecting the crude product to HPLC there was prepared (R)-N-[1-(benzothiazole-2-carbonyl)-butyl]-2-dimethylamino-3-phenylmethanesulfonyl-propionamide (11 mg, 38%).

10 LC/MS retention time 2.98min (TIC), $m/z=488$ (M+H) (determined with method A).

EXAMPLE 19

- (a) (R)-N-[(S)-1-(Benzoxazole-2-carbonyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide

15

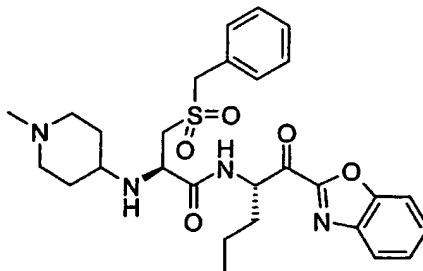


A solution of (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide {0.22mmol, Example 31(a)} in dichloromethane (10mL) was treated with Dess-Martin-periodinane (187mg, 0.44mmol). This mixture was agitated at room temperature overnight then treated with resin-bound thiosulfate (1.47g, 2.2mmol) and stirring was continued for a further 24 hours then the mixture was treated with Silicycle Triamine (611mg, 2.2mmol). After agitating for a further 24 hours the reaction mixture was filtered. The filtrate was evaporated and the residue was subjected to HPLC to give (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide (9mg, 8%). LC/MS

25 retention time 3.0min (TIC), $m/z=528$ (M+H) (determined with method B).

-112-

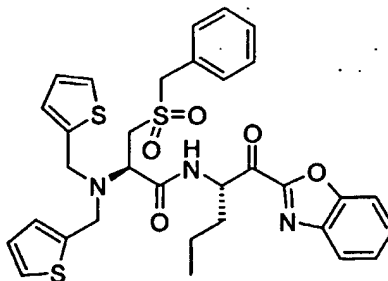
- (b) (R)-N-[(S)-1-(Benzoxazole-2-carbonyl)-butyl]-2-(1-methyl-piperidin-4-ylamino)-3-phenylmethanesulfonyl-propionamide



- 5 By proceeding in a similar manner to Example 19(a) but using (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-(1-methyl-piperidin-4-ylamino)-3-phenylmethanesulfonyl-propionamide {0.22mmol, Example 31(b)} there was prepared (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-(1-methyl-piperidin-4-ylamino)-3-phenylmethanesulfonyl-propionamide (7mg, 6%).
LC/MS retention time 2.7min (TIC), $m/z=541$ (M+H) (determined with method A).

10

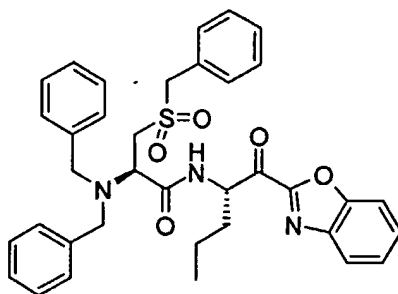
- (c) (R)-N-[(S)-1-(Benzoxazole-2-carbonyl)-butyl]-2-(bis-thiophen-2-ylmethyl-amino)-3-phenylmethanesulfonyl-propionamide



- 15 By proceeding in a similar manner to Example 19(a) but using (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-(bis-thiophen-2-ylmethyl-amino)-3-phenylmethanesulfonyl-propionamide {0.22mmol, Example 31(c)} there was prepared (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-(bis-thiophen-2-ylmethyl-amino)-3-phenylmethanesulfonyl-propionamide (5.3mg, 4%)
LC/MS retention time 3.7min (TIC), $m/z=636$ (M+H) (determined with method A).

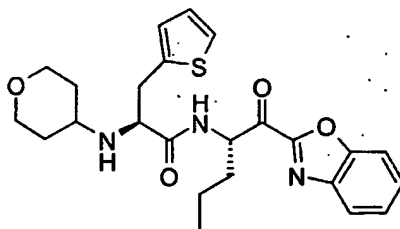
- 20 (d) (R)-N-[(S)-1-(Benzoxazole-2-carbonyl)-butyl]-2-dibenzylamino-3-phenylmethanesulfonyl-propionamide

-113-



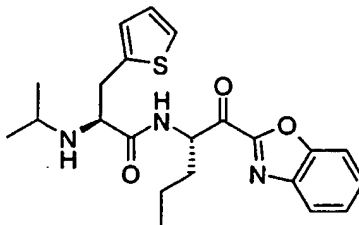
By proceeding in a similar manner to Example 19(a) but using (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-dibenzylamino-3-phenylmethanesulfonyl-propionamide (0.22mmol, Example 31(d)) there was prepared (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-dibenzylamino-3-phenylmethanesulfonyl-propionamide (3.8mg, 3%). LC/MS retention time 4.14min (TIC), $m/z=624$ (M+H) (determined with method B).

(e) (S)-N-[(S)-1-(Benzoxazole-2-carbonyl)-butyl]-2-(tetrahydro-pyran-4-ylamino)-3-thiophen-2-yl-propionamide



By proceeding in a similar manner to Example 19(a) but using (S)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-(tetrahydro-pyran-4-ylamino)-3-thiophen-2-yl-propionamide (0.22mmol, Example 31(e)) there was prepared (S)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-(tetrahydro-pyran-4-ylamino)-3-thiophen-2-yl-propionamide (6.5mg, 6%). LC/MS retention time 2.92min (TIC), $m/z=456$ (M+H) (determined with method B).

(f) (S)-N-[(S)-1-(Benzoxazole-2-carbonyl)-butyl]-2-isopropylamino-3-thiophen-2-yl-propionamide



By proceeding in a similar manner to Example 19(a) but using (S)-N-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-isopropylamino-3-thiophen-2-yl-propionamide (0.22mmol, Example 31(f)),

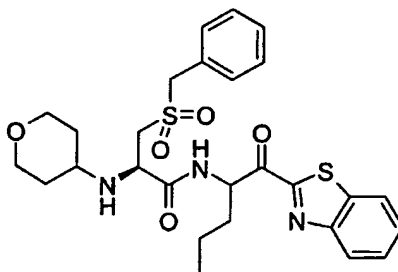
-114-

there was prepared (S)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-isopropylamino-3-thiophen-2-yl-propionamide (10.6mg, 12%). LC/MS retention time 2.99min (TIC), $m/z=414$ (M+H) (determined with method B).

5

EXAMPLE 20

(a) (R)-N-[1-(Benzothiazole-2-carbonyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide



A solution of (R)-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide {0.22mmol, Example 32(a)} in dichloromethane (10mL) was treated with Dess-Martin-periodinane (187mg (0.44mmol)). After stirring at room temperature for 30minutes the reaction mixture was treated with saturated sodium thiosulfate solution (50ml) and saturated sodium bicarbonate solution (50ml). The phases were separated and the aqueous phase extracted with dichloromethane. The combined organic phases were washed with brine, then dried over magnesium sulfate and then evaporated. The residue was subjected to flash chromatography using a silica gel cartridge to give (R)-N-[1-(benzothiazole-2-carbonyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide (46mg, 38%) as mixture of diastereoisomers. The two diastereomers were separated by silica gel column chromatography eluting with 1:1 v/v heptane- ethyl acetate mixture.

20 Diastereoisomer A:

^1H NMR (CDCl_3 , 300MHz): 8.23-8.20 (m, 2H), 8.00 (dd, $J=7\text{Hz}$, 2Hz, 1H), 7.63-7.53 (m, 2H), 7.48-7.40 (m, 5H), 5.80 (m, 1H), 4.38 (d, $J=14\text{Hz}$, 1H), 4.32 (d, $J=14\text{Hz}$, 1H), 3.97-3.90 (m, 2H), 3.80 (dd, $J=9.5\text{Hz}$, 3Hz, 1H), 3.43-3.30 (m, 3H), 3.13 (dd, $J=14.5\text{Hz}$, 9.5Hz, 1H), 2.70 (m, 1H), 2.27 (m, 1H), 2.09 (m, 1H), 1.91-1.76 (m, 3H), 1.52-1.37 (m, 4H), 0.95 (t, $J=7.5\text{Hz}$, 3H).

25 LC/MS $m/z=544$ (M+H)

Diastereoisomer B:

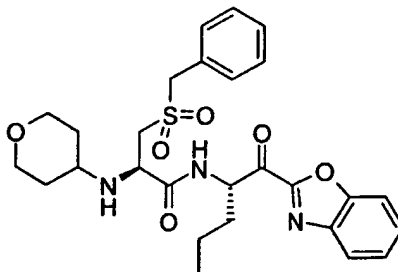
^1H NMR (CDCl_3 , 300MHz): 8.22-8.19 (m, 2H), 8.01-7.98 (m, 1H), 7.63-7.53 (m, 2H), 7.44-7.37 (m, 5H), 5.74 (m, 1H), 4.35-4.31 (m, 2H), 3.99-3.94 (m, 2H), 3.86 (dd $J=9.5\text{Hz}$, 3Hz, 1H), 3.49-3.33 (m, 3H), 3.08 (dd, $J=14.5\text{Hz}$, 9.5Hz), 2.75-2.70 (m, 1H), 2.22 (m, 1H), 2.15-2.06 (m, 1H), 1.91-1.75 (m, 3H), 1.53-1.37 (m, 4H), 0.96 (t, $J=7.5\text{Hz}$, 3H).

30

-115-

LC/MS $m/z=544$ (M+H)

- (b) (R)-N-[(S)-1-(Benzoxazole-2-carbonyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide



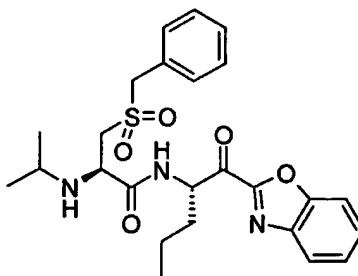
5

By proceeding in a similar manner to Example 20(a) but using (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide {0.22mmol, Example 32(b)} there was prepared (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide (48mg, 41%). ^1H NMR (CDCl_3 , 300MHz): 8.22 (d, $J=8.5\text{Hz}$, 1H), 7.92 (d, $J=8\text{Hz}$, 1H), 7.68 (d, $J=8.5\text{Hz}$, 1H), 7.60-7.40 (m, 7H), 5.68-5.61 (m, 1H), 4.37 (d, $J=14\text{Hz}$, 1H), 4.31 (d, $J=14\text{Hz}$, 1H), 3.97-3.91 (m, 2H), 3.80 (dd, $J=9.5\text{Hz}$, 3H, 1H), 3.43-3.32 (m, 3H), 3.12 (dd, $J=14.5\text{Hz}$, 9.5Hz, 1H), 2.73-2.66 (m, 1H), 2.26 (m, 1H), 2.13-2.05 (m, 1H), 1.89-1.77 (m, 3H), 1.52-1.39 (m, 4H), 0.97 (t, $J=7.5\text{Hz}$, 3H). LC/MS $m/z=528$ (M+H).

15

EXAMPLE 21

- (a) (R)-N-[(S)-1-(Benzoxazole-2-carbonyl)-butyl]-2-isopropylamino-3-phenylmethanesulfonyl-propionamide



A solution of (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-isopropylamino-3-phenylmethanesulfonyl-propionamide {30mg, 0.063mmol, Example 31(g)} in dichloromethane (10mL) was treated with Dess-Martin-periodinane (53mg, 0.126mmol) and this mixture was stirred at room temperature for 1 hour then subjected to The Mettler-Toledo Alexx™ liquid handler work-up as described below:

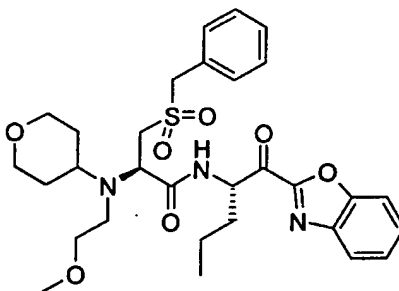
Dichloromethane (15ml) was added to the reaction mixture, followed by a 1:1 mixture (8ml) of saturated sodium thiosulfate solution and saturated sodium bicarbonate solution. The phases were

25

-116-

separated and the organic phase washed with another 5ml of the thiosulfate/bicarbonate solution. The organic phase was then washed with brine and then dried over magnesium sulfate. The crude product was subjected to flash chromatography using a silica gel cartridge to give (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-isopropylamino-3-phenylmethanesulfonyl propionamide (6.2mg, 20%). LC/MS retention time 2.7min (TIC), $m/z=486$ (M+H) (determined with method C).

(b) (R)-N-[(S)-1-(Benzoxazole-2-carbonyl)-butyl]-2-[(2-methoxy-ethyl)-(tetrahydro-pyran-4-yl)-amino]-3-phenylmethanesulfonyl-propionamide

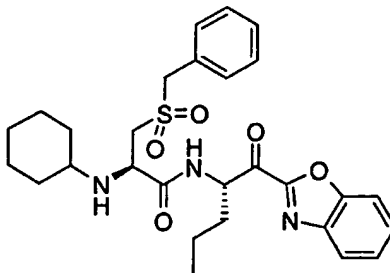


10

By proceeding in a similar manner to Example 21(a) but using (R)-N-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-[(2-methoxy-ethyl)-(tetrahydro-pyran-4-yl)-amino]-3-phenylmethanesulfonyl-propionamide {80mg, 0.136 mmol, Example 32(d)} there was prepared (R)-N-[(S)-1-(Benzoxazole-2-carbonyl)-butyl]-2-[(2-methoxy-ethyl)-(tetrahydro-pyran-4-yl)-amino]-3-phenylmethanesulfonyl-propionamide (7mg, 9%). LC/MS retention time 3.5min (TIC), $m/z=586$ (M+H) (determined with method C).

15

(c) (R)-N-[(S)-1-(Benzoxazole-2-carbonyl)-butyl]-2-cyclohexylamino-3-phenylmethanesulfonyl-propionamide



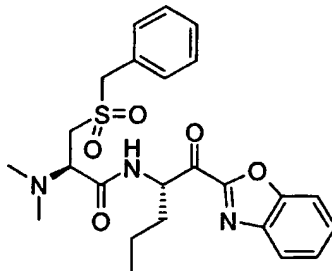
20

By proceeding in a similar manner to Example 21(a) but using (R)-N-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-cyclohexylamino-3-phenylmethanesulfonyl-propionamide {48mg, 0.091mmol, Example 32(e)} there was prepared (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-cyclohexylamino-3-phenylmethanesulfonyl-propionamide (7.9mg, 16%). LC/MS retention time 2.99-3.02min (TIC), $m/z=526$ (M+H) (determined with method C).

25

-117-

- (d) (R)-N-[(S)-1-(Benzoxazole-2-carbonyl)-butyl]-2-dimethylamino-3-phenylmethanesulfonyl-propionamide

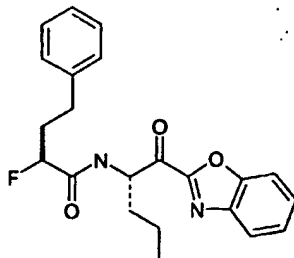


- 5 By proceeding in a similar manner to Example 21(a) but using (R)-N-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-dimethylamino-3-phenylmethanesulfonyl-propionamide {10mg, 0.021mmol, Example 32(f)} there was prepared (R)-N-[(S)-1-(Benzoxazole-2-carbonyl)-butyl]-2-dimethylamino-3-phenylmethanesulfonyl-propionamide (2.5mg, 24%). LC/MS retention time 2.82min (TIC), $m/z=472$ (M+H) (determined with method C).

10

EXAMPLE 22

(1S)-N-[1-(Benzoxazole-2-carbonyl)-butyl]-2-(S)-fluoro-4-phenyl-butyramide



- Step 1. To a mixture of (S)-2-amino-1-benzoxazol-2-yl-pentan-1-ol {0.549 mmol, 121 mg, Reference Example 17(c)}, (S)-2-fluoro-4-phenyl-butyric acid (1.0 eq., 0.549 mmol, 100 mg, Reference Example 9) and N,N-diisopropylethylamine (1.1 eq., 0.604 mmol, 78 mg) in dry dichloromethane (5 mL) under nitrogen was added PyBOP® (1.1 eq., 0.603 mmol, 285 mg). The mixture was stirred at room temperature for 23.5 hr, then concentrated in vacuum. The residue was diluted with ethyl acetate (20 mL) and washed with sodium bicarbonate (30 mL) then water (30 mL). The organic layer was dried (MgSO₄) and concentrated in vacuum. The residue was purified by silica gel column chromatography, eluting with ethyl acetate and heptane (1:2) to afford (S)-N- [(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-fluoro-4-phenyl-butyramide as mixture of diastereoisomers (167.8 mg, 79.5%).

- Step 2. To a solution of (S)-N- [(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-fluoro-4-phenyl-butyramide in dry dichloromethane (5mL) under nitrogen was added a 15% (wt in dichloromethane,

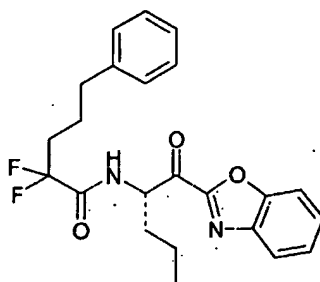
25

-118-

2.0 eq, 0.863 mmol, 2.44 g) of 1,1,1-triacetoxy-1,1-dihydro-1,2-benziodoxol-3(1H)-one (Dess-Martin periodinane). The mixture was stirred at room temperature for 2 hr, then quenched by adding a solution of $\text{Na}_2\text{S}_2\text{O}_3$ (4.0 eq., 1.73 mmol, 273 mg) in saturated Sodium bicarbonate solution (30 ml). The organic layer was dried (MgSO_4) and concentrated in vacuum. The residue was purified over 10 g silica gel, eluting with ethyl acetate and heptane (1:3) to afford (1S)-N-[1-(Benzoxazole-2-carbonyl)-butyl]-2-(S)-fluoro-4-phenyl-butyramide as a light yellow solid (156 mg, 94%). ^1H NMR (CDCl_3) 7.95 (d, $J=7.9$ Hz, 1H), 7.7 (d, $J=8.2$ Hz, 1H), 7.6 (t, $J=7.3$ Hz, 1H), 7.51 (t, $J=7.4$ Hz, 1H), 7.2 (m, 6H), 5.8 (m, 1H), 4.95 (ddd, $J=49.4, 8, 3.5$ Hz, 1H), 2.8 (m, 2H), 2.4 (m, 1H), 2.2 (m, 2H), 1.85 (m, 1H), 1.5 (m, 2H), 1.0 (t, $J=7.3$ Hz, 3H). LC/MS 86% ($M+1$) 383.

EXAMPLE 23

2,2-Difluoro-5-phenyl-pentanoic acid [(S)-1-(benzoxazole-2-carbonyl)-butyl]-amide



Step 1. A solution 2,2-Difluoro-5-phenyl-pentanoic acid (182 mg, 0.85 mmol) in DMF (10 mL) was treated with (S)-2-amino-1-benzoxazol-2-yl-pentan-1-ol (187 mg, 0.85 mmol), HATU (323 mg, 0.85 mmol) and N,N-Diisopropylethylamine (0.162 mL) and stirred at room temperature for 5 ½ hrs. DMF evaporate off, crude taken up in ethyl acetate and washed with 1N HCl, saturated NaHCO_3 and brine. Dried over Na_2SO_4 and evaporated under reduced pressure to give an oil. Purification by column chromatography eluting with 1:1 mixture of ethyl acetate and heptane gave 2,2-Difluoro-5-phenyl-pentanoic acid [(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-amide as orange oil (216 mg). MS 417 (MH^+).

Step 2. A solution of 2,2-Difluoro-5-phenyl-pentanoic acid [(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-amide (216 mg, 0.52 mmol) in dichloromethane (10 mL) was treated with 1,1,1-triacetoxy-1,1-dihydro-1,2-benziodoxol-3(1H)-one (Dess-Martin periodinane) (220 mg, 0.52 mmol) for 1hr at room temperature. The reaction mixture was washed with 0.5 M $\text{Na}_2\text{S}_2\text{O}_3$, saturated NaHCO_3 , and water and dried over Na_2SO_4 . Solvent evaporated under reduced pressure and crude purified by flash chromatography eluting with mixture of ethyl acetate and heptane to give 2,2-Difluoro-5-phenyl-pentanoic acid [(S)-1-(benzoxazole-2-carbonyl)-butyl]-amide as off white solid (90 mg).

-119-

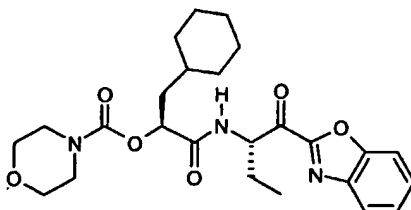
¹H NMR (CDCl₃) 7.93 (d, J=8 Hz, 1H), 7.68 (d, J=8 Hz, 1H), 7.59 (t, J=8 Hz, 1H), 7.49 (t, J=8 Hz, 1H), 7.3-7.11 (m, 5H), 5.72 (m, 1H), 2.67 (t, J=7.5 Hz, 2H), 2.22-2.07 (m, 3H), 1.92-1.77 (m, 3H), 1.55-1.26 (m, 2H), 0.96 (t, J=7.4Hz, 3H).

LC/MS 415(M+1).

5

EXAMPLE 24

(a) Morpholine-4-carboxylic acid (S)-1-[(S)-1-(benzoxazole-2-carbonyl)-propylcarbamoyl]-2-cyclohexyl-ethyl ester



10 Step 1. (S)-3-Cyclohexyl-2-hydroxy-propionic acid (3g, 17.4mmol) was dissolved in methanol (30mL). Trimethylorthoformate (5mL) and p-toluenesulfonic acid monohydrate (100mg) was added. The mixture was stirred at ambient temperature overnight. Water (50mL) was added and stirring was continued for 2h. Methanol was removed under vacuum and the aqueous residue was extracted with ethyl acetate (3x50mL). The combined organic layers were washed with sat. aqueous NaHCO₃ and
15 brine, dried with MgSO₄ and evaporated. (S)-3-Cyclohexyl-2-hydroxy-propionic acid methyl ester was obtained as a colorless liquid (3.1g, 16.7mmol).

Step 2. (S)-3-Cyclohexyl-2-hydroxy-propionic acid methyl ester (1g, 5.37mmol) was dissolved in dichloromethane (20mL). Pyridine (0.57mL, 7mmol) was added and the solution was cooled to 0°C
20 under nitrogen. Trichloromethylchloroformate (0.66mL, 5.5mmol) was added and the mixture was stirred for 30min at room temperature. Morpholine (0.5mL) was added and stirring was continued for 2h. After dilution with ethyl acetate (200mL), the solution was washed with 1N aqueous. HCl and brine, dried with MgSO₄ and evaporated under vacuum. The residue was dissolved in methanol (50mL) and 1N aqueous. NaOH solution (20mL) was added. The mixture was stirred at room
25 temperature for 4h. Methanol was removed under vacuum and the aqueous residue was washed with diethylether. The aqueous layer was acidified with 1N aqueous HCl and extracted with ethyl acetate (3x100mL). The combined organic layers were washed with brine, dried with MgSO₄ and evaporated under vacuum. The crude (S)-morpholine-4-carboxylic acid 1-carboxy-2-cyclohexyl-ethyl ester was used without further purification.

30

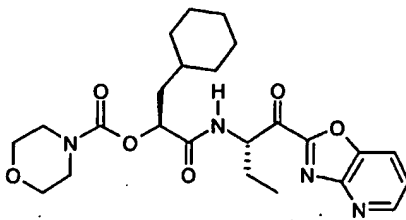
-120-

Step 3. By proceeding in a similar manner to that described in step3 Example 4(a) but using (S)-morpholine-4-carboxylic acid 1-carboxy-2-cyclohexyl-ethyl ester there was prepared morpholine-4-carboxylic acid (S)-1-[(S)-1-(benzooxazole-2-carbonyl)-propylcarbamoyl]-2-cyclohexyl-ethyl ester.

¹H NMR: (DMSO) 8.61 (d, J=6.4Hz, 1H), 7.97 (d, J=8.0Hz, 1H), 7.87 (d, J=8.0Hz, 1H), 7.61 (t, J=8.0Hz, 1H), 7.52 (t, J=8.0Hz, 1H), 5.15-5.09 (m, 1H), 4.91-4.86 (m, 1H), 3.56-3.20 (m, 8H), 2.05-1.93 (m, 1H), 1.79-0.78 (m, 14H), 0.96 (t, J=7.2Hz, 3H). MS: (M+H)⁺ 472.

By proceeding in a similar manner to Example 24(a) there was prepared:

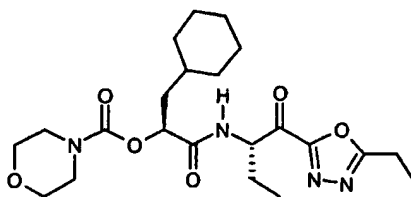
- 10 (b) Morpholine-4-carboxylic acid (S)-2-cyclohexyl-1-[(S)-1-(oxazolo[4,5-b]pyridine-2-carbonyl)-propylcarbamoyl]-ethyl ester



¹H NMR: (DMSO) 8.73-8.69 (m, 2H), 8.38 (d, J=8.0Hz, 1H), 7.67-7.62 (m, 1H), 5.08-5.02 (m, 1H), 4.88-4.83 (m, 1H), 3.57-3.20 (m, 8H), 2.07-1.95 (m, 1H), 1.79-0.75 (m, 14H), 0.97 (t, J=7.2Hz, 3H).

15 MS: (M+H)⁺ 473;

- (c) Morpholine-4-carboxylic acid (S)-2-cyclohexyl-1-[(S)-1-(5-ethyl-[1,3,4]oxadiazole-2-carbonyl)-propylcarbamoyl]-ethyl ester

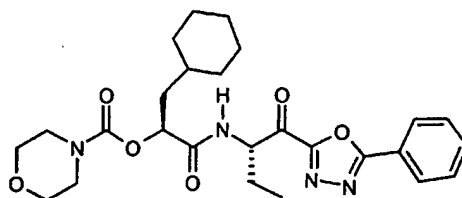


20 ¹H NMR: (DMSO) 8.62 (d, J=4.8Hz, 1H), 4.94-4.84 (m, 2H), 3.57-3.20 (m, 8H), 2.95 (q, J=7.2Hz, 2H), 1.98-1.87 (m, 1H), 1.74-0.82 (m, 14H), 1.29 (t, J=7.2Hz, 3H), 0.93 (t, J=7.2Hz, 3H). MS: (M+H)⁺ 451;

- (d) Morpholine-4-carboxylic acid (S)-2-cyclohexyl-1-[(S)-1-(5-phenyl-[1,3,4]oxadiazole-2-carbonyl)-propylcarbamoyl]-ethyl ester

25

-121-

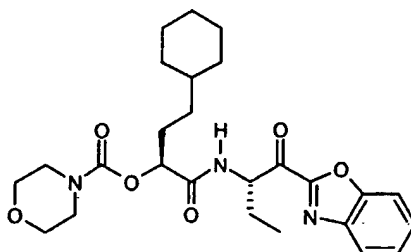


¹H NMR: (DMSO) 8.69 (d, J=6.0Hz, 1H), 8.07 (d, J=8Hz, 2H), 7.70-7.59 (m, 3H), 4.99-4.92 (m, 1H), 4.88-4.83 (m, 1H), 3.57-3.20 (m, 8H), 2.03-1.92 (m, 1H), 1.77-0.77 (m, 14H), 0.96 (t, J=7.2Hz, 3H).

MS: (M+H)⁺ 499;

5

(e) Morpholine-4-carboxylic acid (S)-1-[(S)-1-(benzooxazole-2-carbonyl)-propylcarbamoyl]-3-cyclohexyl-propyl ester

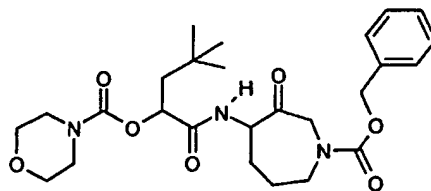


¹H NMR: (DMSO) 8.60 (d, J=6.8Hz, 1H), 7.97 (d, J=8.0Hz, 1H), 7.87 (d, J=8.0Hz, 1H), 7.61 (t, J=8.0Hz, 1H), 7.52 (t, J=8.0Hz, 1H), 5.13-5.06 (m, 1H), 4.81-4.76 (m, 1H), 3.56-3.21 (m, 8H), 2.05-1.93 (m, 1H), 1.79-1.46 (m, 8H), 1.19-0.90 (m, 6H), 0.96 (t, J=7.2Hz, 3H), 0.77-0.62 (m, 2H). MS: (M+H)⁺ 486;

10

EXAMPLE 25

15 4-[4,4-Dimethyl-2-(morpholine-4-carboxyloxy)-pentanoylamino]-3-oxo-azepane-1-carboxylic acid benzyl ester



20

Sodium hydride (60% in mineral oil, 10g, 250mmol) was suspended in dry DMF. Allyl-carbamic acid benzyl ester (19.1g, 100mmol) was added dropwise at ambient temperature. After stirring for 5min, 5-bromo-1-pentene (25g, 168mmol) was added dropwise. Stirring was continued at 50°C for 1h. The reaction was quenched with water and then partitioned between diethylether and water. The ether layer was washed with water and brine, dried with MgSO₄ and evaporated under vacuum. Flash chromatography (ethyl acetate/hexane 1:9) gave 15.5g allyl-pent-4-enyl-carbamic acid benzyl ester.

-122-

Allyl-pent-4-enyl-carbamic acid benzyl ester (15.5g, 59.8mmol) was dissolved in dichloromethane and bis(tricyclohexylphosphine)benzylidene ruthenium(IV) dichloride (1g) was added. The mixture was refluxed under a nitrogen atmosphere until TLC analysis showed complete reaction. The solvent was evaporated under vacuum and the residue was purified by flash chromatography (ethyl acetate/hexane 1:9). Yield: 7.8g 2,3,4,7-Tetrahydro-azepine-1-carboxylic acid benzyl ester.

To a solution of 2,3,4,7-tetrahydro-azepine-1-carboxylic acid benzyl ester (4.5g, 19.45mmol) in dichloromethane (50mL) was added m-chloroperbenzoic acid (60mmol). The mixture was stirred at ambient temperature for 16h. Sat aqueous K_2CO_3 solution was added and the mixture was extracted with dichloromethane. The combined organic layers were washed with sat. aqueous $NaHCO_3$ and brine, dried with $MgSO_4$ and evaporated under vacuum. The crude epoxide was dissolved in a 8:1 methanol/water mixture (100mL). Ammonium chloride (3.2g, 60mmol) and sodium azide (3.9g, 60mmol) was added and the mixture was heated at 60°C for 48h. Most of the solvent was removed under vacuum. The residue was extracted with ethyl acetate. The combined organic layers were washed with sat. aqueous $NaHCO_3$ (200mL) and brine (200mL), dried with $MgSO_4$ and evaporated under vacuum. Flash chromatography of the residue (hexane/ethyl acetate 3:1) gave 3.3g of 4-azido-3-hydroxy-azepane-1-carboxylic acid benzyl ester.

To a solution of 4-azido-3-hydroxy-azepane-1-carboxylic acid benzyl ester (3.3g, 11.37mmol) in methanol (50mL) was added triethylamine (5mL) and 1,3-propanedithiol (3.42mL, 35mmol). The mixture was stirred at ambient temperature until TLC analysis showed complete consumption of the starting material. A white precipitate was removed by filtration and the filtrate was evaporated to dryness. The residue was triturated with a 1:1 hexane/diethylether mixture to remove excess dithiol and dried under vacuum.

Crude 4-amino-3-hydroxy-azepane-1-carboxylic acid benzyl ester (150mg, 0.57mmol), morpholine-4-carboxylic acid 1-carboxy-3,3-dimethyl-butyl ester (120mg, 0.46mmol), EDC (400mg, 2.1mmol), and HOBt (400mg, 2.5mmol) were combined. Dichloromethane (5mL) was added and then 4-methylmorpholine (0.5mL). The mixture was stirred at ambient temperature for 2h. After dilution with ethyl acetate (100mL) the solution was washed with 1N HCl, sat. aqueous $NaHCO_3$ and brine, dried with $MgSO_4$ and evaporated under vacuum. The residue was dissolved in DMSO (5mL). Triethylamine (0.3mL) and then SO_3 pyridine complex (150mg) were added and the mixture was stirred at ambient temperature for 2h. After dilution with ethyl acetate (100mL), the solution was washed with water (50mL) and brine, dried with $MgSO_4$ and evaporated under vacuum. The residue was purified by flash chromatography on silica gel and gave 4-[4,4-Dimethyl-2-(morpholine-4-

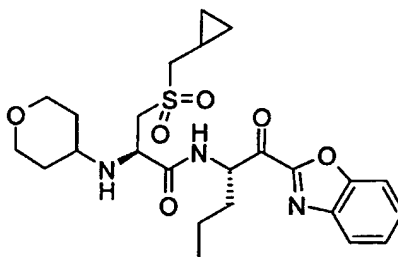
-123-

carbonyloxy)-pentanoylamino]-3-oxo-azepane-1-carboxylic acid benzyl ester (95mg, 0.189mmol) as a white solid.

2:1 mixture of diastereomers, ¹H NMR: (DMSO) 8.14-8.08 (m, 1H), 7.40-7.25 (m, 5H), 5.18-4.89 (m, 3H), 4.51-4.33 (m, 2H), 4.01-3.76 (m, 2H), 3.60-3.25 (m, 8H), 2.95-2.79 (m, 1H), 1.84-1.54 (m, 6H), 0.92/0.91 (s, 9H). MS: (M+H)⁺ 504. LC/MS m/z=474(M+H)

EXAMPLE 26

(a) (R)-N-[(S)-1-(Benzoxazole-2-carbonyl)-butyl]-3-cyclopropylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide



10

Step 1. (R)-2-Amino-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-cyclopropylmethanesulfonyl-propionamide (90mg, 0.22mmol, Reference Example 11(f)) was dissolved in 5% acetic acid in acetonitrile (10ml). Tetrahydro-4H-pyran-4-one (110mg, 1.1mmol) was added, followed by (polystyrylmethyl)trimethylammonium cyanoborohydride (107mg, 1.1mmol). The resulting reaction mixture was stirred for four hours and then filtered under suction. The solvents were evaporated under high vacuum. The residue was dissolved in 5ml dichloromethane, Silicycle Triamine (940mg, 2.2mmol) was added and the reaction mixture stirred for four hours. It was filtered under suction and the filtrate concentrated under reduced pressure to give (R)-N-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-cyclopropylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide (89mg, 0.18mmol, 82%).

20

Step 2. (R)-N-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-cyclopropylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide (89mg, 0.18mmol) was dissolved in 10ml dichloromethane. The Dess-Martin-periodinane (153mg, 0.36mmol) was added and the resulting reaction mixture stirred for two hours. The reaction mixture was poured into a 1/1-mixture of saturated sodium bicarbonate solution and saturated sodium thiosulfate solution. The aqueous phase was extracted with dichloromethane. The combined organic phases were washed with saturated sodium bicarbonate solution and brine. The organic phase was dried with magnesium sulfate and the dichloromethane evaporated under reduced pressure. The crude product was purified via flash chromatography (heptane/ethyl acetate 1/1 to elute) to give (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-3-cyclopropylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide (24mg,

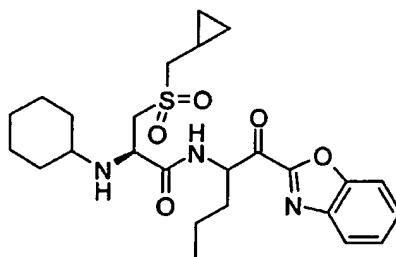
30

-124-

0.049mmol, 27%). ¹H NMR (CDCl₃, 300MHz): 8.29 (d, J=8.5Hz, 1H), 7.93 (d, J=8Hz, 1H), 7.68 (d, J=8Hz, 1H), 7.59-7.46 (m, 2H), 5.67 (m, 1H), 3.99-3.93 (m, 2H), 3.84 (dd, J=9.5Hz, 2.5Hz, 1H), 3.56 (dd, J=14.5Hz, 2.5Hz, 1H), 3.42-3.33 (m, 2H), 3.24 (dd, J=14.5Hz, 9.5Hz, 1H), 3.02-2.99 (m, 2H), 2.78-2.71 (m, 1H), 2.13-2.07 (m, 1H), 1.95-1.78 (m, 3H), 1.55-1.41 (m, 5H), 1.23-1.16 (m, 1H), 1.00 (t, J=7.5Hz, 3H), 0.81-0.74 (m, 2H), 0.48-0.43 (m, 2H).

LC/MS m/z=492 (M+H)

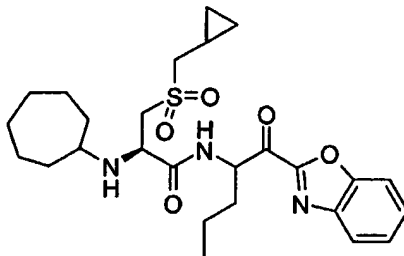
(b) (R)-N-[1-(benzoxazole-2-carbonyl)-butyl]-2-cyclohexylamino-3-cyclopropylmethanesulfonyl-propionamide



By proceeding in a similar manner to Example 26(a) but using cyclohexanone there was prepared

(R)-N-[1-(benzoxazole-2-carbonyl)-butyl]-2-cyclohexylamino-3-cyclopropylmethanesulfonyl-propionamide (predominantly as one diastereomer). ¹H NMR (CDCl₃, 300MHz): 8.37 (d, J=8.5Hz, 1H); 7.92 (d, J=8Hz, 1H), 7.67 (d, J=8Hz, 1H), 7.59-7.36 (m, 2H), 5.65 (m, 1H), 3.79 (dd, J=9.5Hz, 2.5Hz, 1H), 3.54 (dd, J=14.25Hz, 2.5Hz, 1H), 3.24 (dd, J=14.25Hz, 9.5Hz, 1H), 3.02-2.95 (m, 2H), 2.49 (m, 1H), 2.12-2.07 (m, 1H), 1.96-1.17 (m, 15H), 0.98 (t, J=7Hz, 3H), 0.80-0.72 (m, 2H), 0.48-0.43 (m, 2H). LC/MS m/z=490 (M+H)

(c) (R)-N-[1-(Benzoxazole-2-carbonyl)-butyl]-2-cycloheptylamino-3-cyclopropylmethanesulfonyl-propionamide



By proceeding in a similar manner to Example 26(a) but using cycloheptanone there was prepared

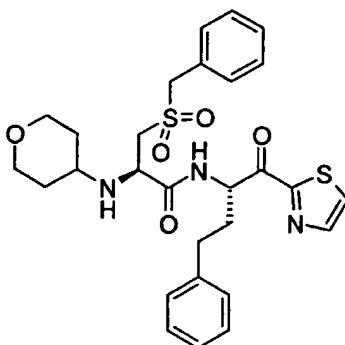
(R)-N-[1-(benzoxazole-2-carbonyl)-butyl]-2-cycloheptylamino-3-cyclopropylmethanesulfonyl-propionamide ¹H NMR (CDCl₃, 300MHz): [8.36 (d, J=8.5Hz), 8.28 (d, J=8.5Hz), 1H], [8.05 (dd, J=8Hz, 1Hz), 7.97 (dd, J=8.5Hz, 1.5Hz), 1H], [7.92 (d, J=8.5Hz), 7.67 (d, J=8Hz), 1H], 7.59-7.48 (m, 1H), [7.44 (ddd, J=8Hz, 7.5Hz, 1Hz), 7.19 (ddd, J=8Hz, 7.5Hz, 1Hz), 1H], [5.65 (m), 5.62 (m), 1H],

-125-

[3.82 (dd, J=10Hz, 3Hz), 3.75 (dd, J=9Hz, 3Hz), 1H], [3.55 (dd, J=14.5Hz, 3Hz), 3.49 (dd, J=14.5Hz, 3Hz), 1H], 3.27 (dd, J=14.5Hz, 9Hz, 1H), 3.03-2.96 (m, 2H), 2.72 (m, 1H), 2.14-2.05 (m, 1H), 1.91-1.39 (m, 16H), 1.23-1.17 (m, 1H), [0.99 (t, J=7.25Hz), 0.98 (t, J=7.25Hz), 1H], 0.79-0.7 (m, 2H), 0.48-0.44 (m, 2H). LC/MS m/z=504 (M+H).

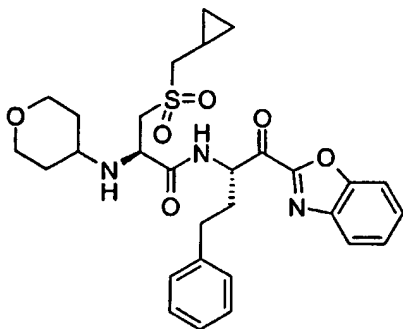
5

- (d) (R)-3-Phenylmethanesulfonyl-N-[(S)-3-phenyl-1-(thiazole-2-carbonyl)-propyl]-2-(tetrahydro-pyran-4-ylamino)-propionamide



By proceeding in a similar manner to Example 26(a) but using (R)-2-Amino-N-[(S)-1-(hydroxy-
 10: thiazol-2-yl-methyl)-3-phenyl-propyl]-3-phenylmethanesulfonyl-propionamide {Reference Example
 11(k)} there was prepared (R)-3-phenylmethanesulfonyl-N-[(S)-3-phenyl-1-(thiazole-2-carbonyl)-
 propyl]-2-(tetrahydro-pyran-4-ylamino)-propionamide. ¹H NMR (CDCl₃, 300MHz): 8.27 (d, J=9Hz,
 1H), 8.06 (d, J=3Hz, 1H), 7.73 (d, J=3Hz, 1H), 7.47-7.39 (m, 5H), 7.25-7.11 (m, 5H), 5.72 (m, 1H),
 4.36 (d, J=14Hz, 1H), 4.31 (d, J=14Hz, 1H), 3.97-3.90 (m, 2H), 3.76 (dd, J=9.5Hz, 3Hz, 1H), 3.40-
 15 3.31 (m, 3H), 3.01 (dd, J=14.5Hz, 9.5Hz, 1H), 2.76-2.62 (m, 3H), 2.51-2.40 (m, 1H), 2.22-2.09 (m,
 1H), 1.87-1.75 (m, 2H), 1.53-1.38 (m, 3H)
 LC/MS m/z=556 (M+H);

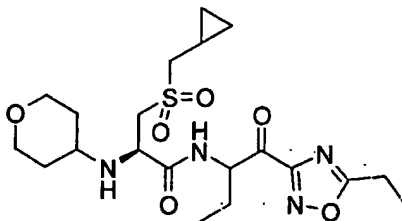
- (e) (R)-N-[(S)-1-(Benzoxazole-2-carbonyl)-3-phenyl-propyl]-3-cyclopropylmethanesulfonyl-2-(
 20 tetrahydro-pyran-4-ylamino)-propionamide



-126-

By proceeding in a similar manner to Example 26(a) but using (R)-2-amino-N-[(S)-1-(hydroxy-thiazol-2-yl-methyl)-3-phenyl-propyl]-3-phenylmethanesulfonyl-propionamide {Reference Example 11(j)} there was prepared (R)-N-[(S)-1-(Benzoxazole-2-carbonyl)-3-phenyl-propyl]-3-cyclopropylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide. ¹H NMR (CDCl₃, 300MHz): 8.36 (d, J=8.5Hz, 1H), 7.92 (d, J=8Hz, 1H), 7.67 (d, J=8Hz, 1H), 7.60-7.46 (m, 2H), 7.25-7.16 (m, 5H), 5.72 (m, 1H), 3.99-3.93 (m, 2H), 3.81 (dd, J=9.5Hz, 3Hz, 1H), 3.52 (dd, J=14Hz, 3Hz, 1H), 3.41-3.33 (m, 2H), 3.15 (dd, J=14Hz, 9.5Hz, 1H), 3.01-2.70 (m, 2H), 2.81-2.70 (m, 3H), 2.53 (m, 1H), 2.27-2.23 (m, 1H), 1.94-1.44 (m, 5H), 1.22-1.17 (m, 1H), 0.80-0.74 (m, 2H), 0.47-0.42 (m, 2H). LC/MS m/z=554 (M+H);

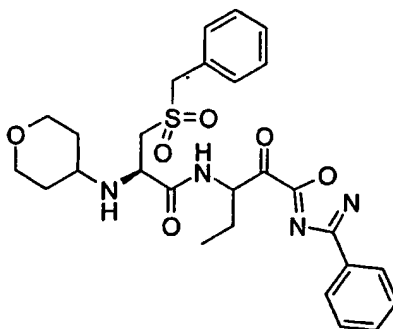
(f) (R)-3-Cyclopropylmethanesulfonyl-N-[1-(5-ethyl-1,2,4-oxadiazole-3-carbonyl)-propyl]-2-(tetrahydro-pyran-4-ylamino)-propionamide



By proceeding in a similar manner to Example 26(a) but using (R)-2-Amino-3-cyclopropylmethanesulfonyl-N-[(S)-1-[(5-ethyl-1,2,4-oxadiazol-3-yl)-hydroxy-methyl]-propyl]-propionamide {Reference Example 11(h)} there was prepared (R)-3-cyclopropylmethanesulfonyl-N-[1-(5-ethyl-1,2,4-oxadiazole-3-carbonyl)-propyl]-2-(tetrahydro-pyran-4-ylamino)-propionamide. ¹H NMR (CDCl₃, 300MHz): [8.28 (d, J=8.5Hz), 8.15 (d, J=8Hz), 1H], [5.40 (m), 5.33 (m), 1H], 3.99-3.95 (m, 2H), [3.90 (dd, J=10Hz, 3Hz), 3.84 (dd, J=9.5Hz, 3Hz), 1H], [3.55 (dd, J=14Hz, 3Hz), 3.47 (dd, J=14Hz, 11Hz), 1H], 3.45-3.33 (m, 2H), 3.23 (dd, 14Hz, 10Hz, 1H), 3.07-2.94 (m, 4H), 2.82-2.71 (m, 1H), 2.19-2.08 (m, 1H), 1.95-1.77 (m, 5H), 1.58-1.43 (m, 1H), 1.45 (t, J=7.5Hz, 3H), 1.23-1.14 (m, 1H), [1.00 (t, J=7.5Hz), 0.97 (t, J=7.5Hz), 3H], 0.81-0.73 (m, 2H), 0.48-0.41 (m, 2H). LC/MS m/z=457 (M+H);

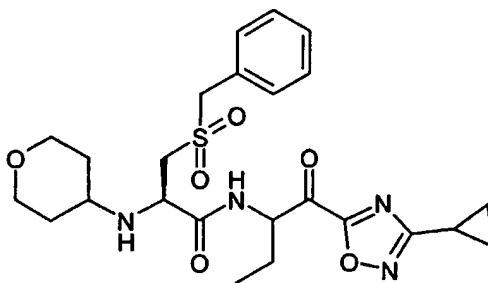
(g) (R)-3-Phenylmethanesulfonyl-N-[1-(3-phenyl-1,2,4-oxadiazole-5-carbonyl)-propyl]-2-(tetrahydro-pyran-4-ylamino)-propionamide

-127-



By proceeding in a similar manner to Example 26(a) but using (R)-2-Amino-N-{1-[hydroxy-(3-phenyl-1,2,4-oxadiazol-5-yl)-methyl]-propyl}-3-phenylmethanesulfonyl-propionamide {Reference Example 11(g)} there was prepared (R)-3-phenylmethanesulfonyl-N-[1-(3-phenyl-1,2,4-oxadiazole-5-carbonyl)-propyl]-2-(tetrahydro-pyran-4-ylamino)-propionamide. ¹H NMR (CDCl₃, 300MHz): [8.15 (d, J=8Hz), 8.14 (d, J=8Hz), 1H], 7.61-7.39 (m, 10H), [5.46 (m), 5.40 (m), 1H], 4.34-4.28 (m, 2H), 4.09-3.93 (m, 2H), [3.87 (dd, J=9.5Hz, 3Hz), 3.81 (dd, J=9.5Hz, 3Hz), 1H], 3.41-3.32 (m, 3H), [3.16 (dd, J=13.5Hz, 10Hz), 3.11 (dd, J=14Hz, 9.5Hz), 1H], 2.75-2.68 (m, 1H), 2.23-2.13 (m, 1H), 1.96-1.43 (m, 6H), 1.06-0.99 (m, 3H), LC/MS m/z=541 (M+H).

(h) (R)-N-[1-(3-Cyclopropyl-1,2,4-oxadiazole-5-carbonyl)-propyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide



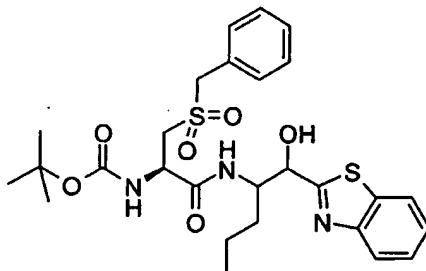
By proceeding in a similar manner to Example 26(a) but using (R)-2-Amino-3-phenylmethanesulfonyl-N-[(S)-1-[(3-cyclopropyl-1,2,4-oxadiazol-5-yl)-hydroxy-methyl]-propyl]-propionamide {Reference Example 11(l)} there was prepared (R)-N-[1-(3-cyclopropyl-1,2,4-oxadiazole-5-carbonyl)-propyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide. ¹H NMR (CDCl₃, 300MHz): [8.19 (d, J=8.5Hz), 8.11 (d, J=7.5Hz), 1H], 7.46-7.40 (m, 5H), [5.33 (m), 5.27 (m), 1H], 4.55-4.35 (m, 2H), 3.99-3.95 (m, 2H), [3.88 (dd, J=10Hz, 3Hz), 3.83 (dd, J=9.5Hz, 3Hz), 1H], 3.44-3.34 (m, 3H), 3.18-3.07 (m, 1H), 2.78-2.67 (m, 1H), 2.24-2.17 (m, 1H), 2.15-2.08 (m, 1H), 1.89-1.72 (m, 3H), 1.55-1.43 (m, 2H), 1.20-1.11 (m, 4H), [0.98 (t, J=7.5Hz), 0.97 (t, J=7.5Hz), 3H].

LC/MS m/z=505 (M+H).

-128-

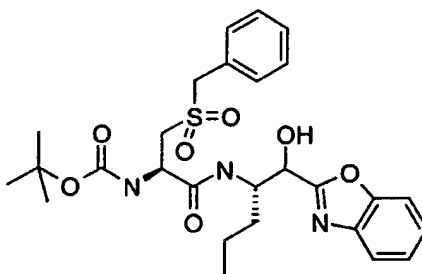
EXAMPLE 27

- (a) {{(R)-1-[1-(Benzothiazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}}-carbamic acid tert-butyl ester



5 N-cyclohexylcarbodiimide, N'-methyl polystyrene (1.74g, 3.4mmol) suspended in a mixture of dichloromethane (10ml) and dimethylformamide (2mL) was treated with hydroxybenzotriazole (391mg, 2.89mmol) and L-N-boc-benzylsulfonylalanine (876mg, 2.55mmol). This mixture was stirred at room temperature for 30 minutes, then treated with 2-amino-1-benzothiazol-2-yl-pentan-1-ol {400mg, 1.7mmol, Reference Example 17(d)} and after stirring for a further 2 hours the mixture was
10 then treated with Silicycle-Triamine (2.36g, 8.5mmol). The reaction mixture was stirred for 2 hours and then filtered. The filtrate was evaporated to give the title compound (888mg, 93%). LC/MS m/z=562.

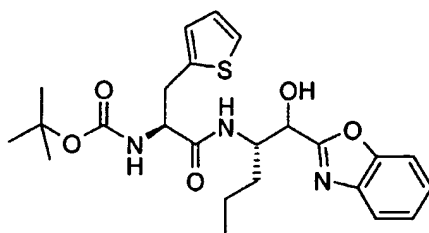
- (b) {{(R)-1-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}}-carbamic acid tert-butyl ester



By proceeding in a manner similar to Example 27(a) above but using L-N-boc-benzylsulfonylalanine (876mg, 2.55mmol) and (2S)-2-amino-1-benzoxazol-2-yl-pentan-1-ol {374mg, 1.7mmol, Reference Example 17(c)} there was prepared {{(R)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}}-carbamic acid tert-butyl ester (908mg, 98%).
20

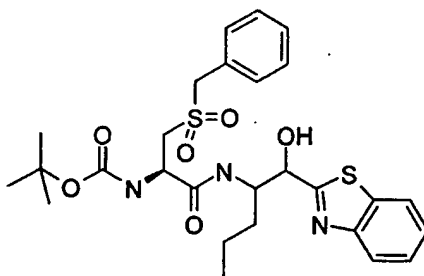
- (c) {{(S)-1-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-thiophen-2-yl-ethyl}}-carbamic acid tert-butyl ester

-129-



By proceeding in a manner similar to Example 27(a) above but using Resin-bound diimide (1.76g, 3.4mmol) suspended in dichloromethane (10mL), hydroxybenzotriazole (391mg, 2.89mmol), (2S)- 2-tert-butoxycarbonylamino-3-thiophen-2-yl- propionic acid (692mg, 2.55mmol), (2S)-2-amino-1-benzoxazol-2-yl-pentan-1-ol {374mg, 1.7mmol, Reference Example 17(c)} and Silicycle-Triamine (2.36g, 8.5mmol) there was prepared {(S)-1-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-thiophen-2-yl-ethyl}-carbamic acid tert-butyl ester (790mg (1.67mmol, 98%). LC/MS:m/z=562 (M+H).

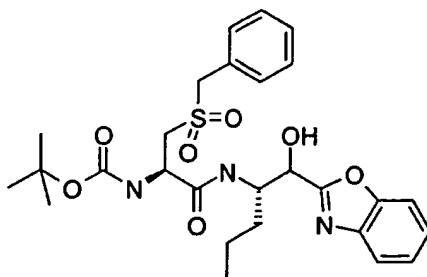
- 10 (d) {(R)-1-[1-(Benzothiazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester



By proceeding in a manner similar to Example 27(a) above but using Resin-bound diimide (741mg, 1.26mmol), hydroxybenzotriazole (144mg, 1.07mmol), L-N-boc-benzylsulfonylalanine (326mg, 0.95mmol), 2-amino-1-benzothiazol-2-yl-pentan-1-ol {150mg, 0.63mmol, Reference Example 17(d)} and Silicycle-Triamine (2.36g, 8.5mmol) there was prepared {(R)-1-[1-(benzothiazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester, LC/MS m/z=562 (M+H), which was used without further purification

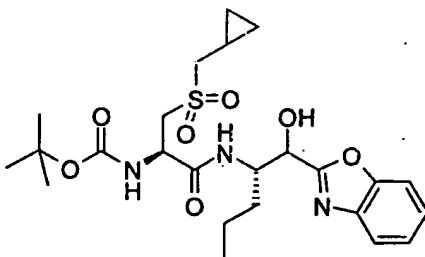
- 20 (e) {(R)-1-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester

-130-



By proceeding in a manner similar to Example 27(a) above but using Resin-bound diimide (1.76g, 3.4mmol), hydroxybenzotriazole (391mg, 2.89mmol), L-N-boc-benzylsulfonylalanine (876mg, 2.55mmol), (2S)-2-amino-1-benzoxazol-2-yl-pentan-1-ol (374mg, 1.7mmol, Reference Example 17(c)) and Silicycle-Triamine (2.36g, 8.5mmol) there was prepared {(R)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester, LC/MS m/z=546 (M+H), 490 (M=H-butene), which was used directly in the next reaction.

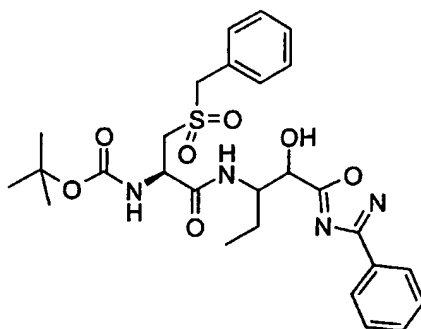
(f) {(R)-1-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-cyclopropylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester



By proceeding in a manner similar to Example 27(a) above but using a suspension of Resin-bound diimide (1.07g, 1.82mmol) in dichloromethane (20ml), hydroxybenzotriazole (209mg, 1.55mmol) and (R)-2-tert-butoxycarbonylamino-3-cyclopropylmethanesulfonyl-propionic acid (420mg, 1.365mmol, Reference Example 22), (S)-2-amino-1-benzoxazol-2-yl-pentan-1-ol (200mg 0.91mmol, Reference Example 17(c)) and Silicycle-Triamine (2.8g, 9.1mmol) there was prepared {(R)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-cyclopropylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester (450mg, 97%). LC/MS m/z=532(M+Na), 510 (M+H), 454 (M+H-isobutene).

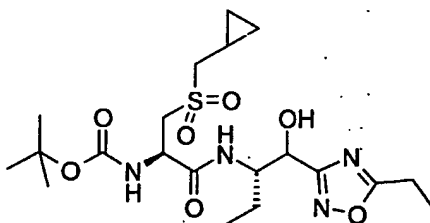
(g) {(R)-1-[(S)-1-(Hydroxy-(3-phenyl-1,2,4-oxadiazol-5-yl)-methyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester

-131-



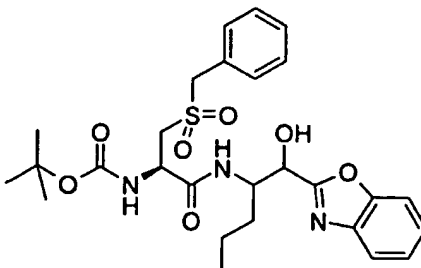
By proceeding in a manner similar to Example 27(f) above but using L-N-boc-benzylsulfonylalanine and (R)-2-tert-butoxycarbonylamino-3-phenylmethanesulfonyl-propionic acid and (S)-2-amino-1-(3-phenyl-[1,2,4]oxadiazol-5-yl)-butan-1-ol (Reference Example 21) there was prepared (R)-1-{1-
 5 [hydroxy-(3-phenyl-1,2,4-oxadiazol-5-yl)-methyl]-propylcarbamoyl}-2-phenylmethanesulfonyl-ethyl-carbamic acid tert-butyl ester. LC/MS m/z=545(M+Na), 467 (M+H-isobutene), 423 (M+H-Boc).

- (i) ((R)-2-Cyclopropylmethanesulfonyl-1-{(S)-1-[(5-ethyl-1,2,4-oxadiazol-3-yl)-hydroxy-methyl]-propylcarbamoyl}-ethyl)-carbamic acid tert-butyl ester



By proceeding in a manner similar to Example 27(f) above but using 2-amino-1-(5-ethyl-[1,2,4]-oxadiazol-3-yl)-butan-1-ol (Reference Example 23) there was prepared ((R)-2-
 10 cyclopropylmethanesulfonyl-1-{(S)-1-[(5-ethyl-1,2,4-oxadiazol-3-yl)-hydroxy-methyl]-propylcarbamoyl}-ethyl)-carbamic acid tert-butyl ester. LC/MS m/z=497(M+Na), 419 (M+H-isobutene), 375 (M+H-Boc).

- (j) {(R)-1-[1-(Benzoxazol-2-yl)-hydroxy-methyl]-butylcarbamoyl}-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester

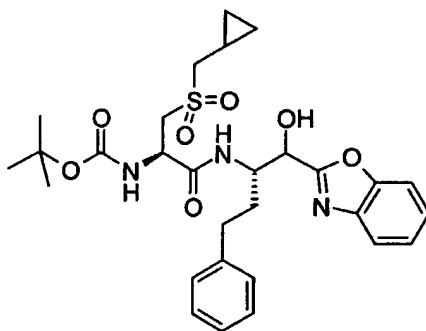


-132-

By proceeding in a manner similar to Example 27(f) above but using L-N-boc-benzylsulfonylalanine and (S)-2-amino-1-benzoxazol-2-yl-pentan-1-ol {Reference Example 17(c)} there was prepared {(R)-1-[1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester. LC/MS $m/z=546(M+H)$, 490 ($M+H$ -isobutene).

5

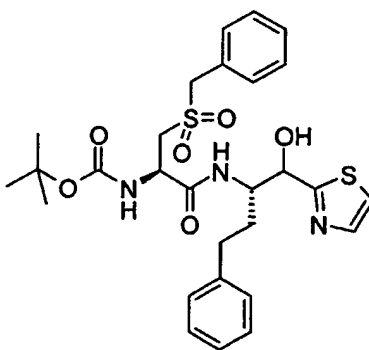
- (k) {(R)-1-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-3-phenyl-propylcarbamoyl]-2-cyclopropylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester



- By proceeding in a manner similar to Example 27(f) above but using (2S)-2-amino-4-phenyl-1-benzoxazol-2-yl-butan-1-ol there was prepared {(R)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-3-phenyl-propylcarbamoyl]-2-cyclopropylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester. LC/MS $m/z=572(M+H)$, 516 ($M+H$ -isobutene).

- (l) {(R)-1-[(S)-1-(Hydroxy-thiazol-2-yl-methyl)-3-phenyl-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester

15

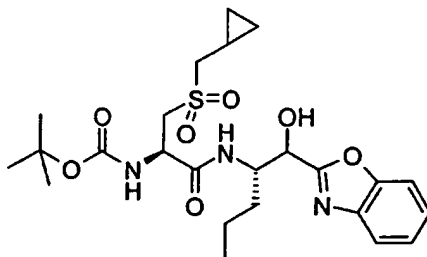


- By proceeding in a manner similar to Example 27(f) above but using L-N-boc-benzylsulfonylalanine and (2S)-2-amino-4-phenyl-1-thiazol-2-yl-butan-1-ol (Reference Example 13) there was prepared {(R)-1-[(S)-1-(hydroxy-thiazol-2-yl-methyl)-3-phenyl-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester. LC/MS $m/z=574(M+H)$.

20

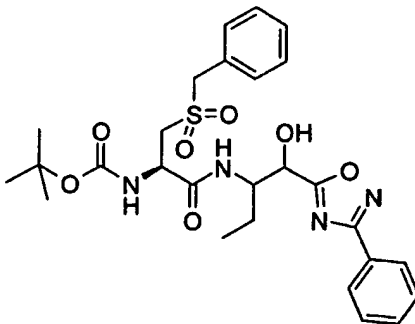
-133-

- (m) {(R)-1-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-cyclopropylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester



By proceeding in a manner similar to Example 27(f) above but using *N*-Cyclohexylcarbodiimide, *N*'-methyl polystyrene (1.07g, 1.82mmol) suspended in dichloromethane (20mL), hydroxybenzotriazole (209mg, 1.55mmol), (R)-2-tert-butoxycarbonylamino-3-cyclopropylmethanesulfonyl-propionic acid (420mg, 1.365mmol, Reference Example 22), (S)-2-amino-1-benzoxazol-2-yl-pentan-1-ol {200mg 0.91mmol, Reference Example 17(c)} and Silicycle-Triamine (2.8g, 9.1mmol) there was prepared {(R)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-cyclopropylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester (450mg, 0.88mmol, 97%). LC/MS m/z =532(M+Na), 510 (M+H), 454 (M+H-isobutene).

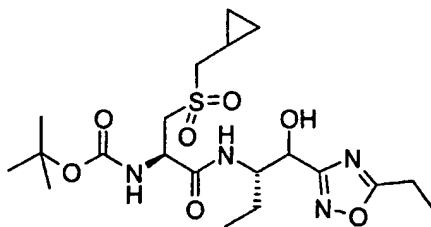
- (n) (R)-1-[1-[Hydroxy-(3-phenyl-1,2,4-oxadiazol-5-yl)-methyl]-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester



By proceeding in a manner similar to Example 27(m) above but using *L*-N-boc-benzylsulfonylalanine and (S)-2-amino-1-(3-phenyl-[1,2,4]oxadiazol-5-yl)-butan-1-ol (Reference Example 21) there was prepared (R)-1-[1-[hydroxy-(3-phenyl-1,2,4-oxadiazol-5-yl)-methyl]-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester. LC/MS m/z =545(M+Na), 467 (M+H-isobutene), 423 (M+H-Boc).

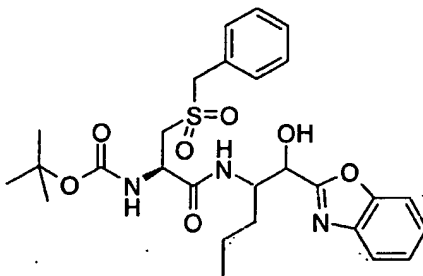
- (o) {(R)-2-Cyclopropylmethanesulfonyl-1-[(S)-1-[(5-ethyl-1,2,4-oxadiazol-3-yl)-hydroxy-methyl]-propylcarbamoyl]-ethyl}-carbamic acid tert-butyl ester

-134-



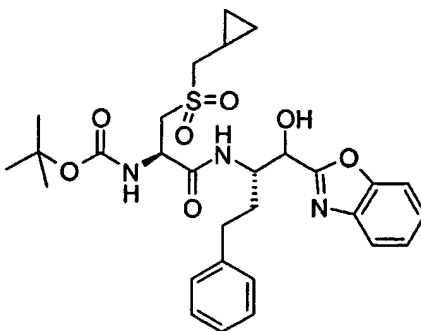
By proceeding in a manner similar to Example 27(m) above but using (S)-2-amino-1-(5-ethyl-[1,2,4]oxadiazol-3-yl)-butan-1-ol there was prepared ((R)-2-Cyclopropylmethanesulfonyl-1-[(S)-1-[(5-ethyl-1,2,4-oxadiazol-3-yl)-hydroxy-methyl]-propylcarbamoyl]-ethyl)-carbamic acid tert-butyl ester. LC/MS $m/z=497(M+Na)$, 419 (M+H-isobutene), 375 (M+H-Boc)

(p) {(R)-1-[1-(Benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester



By proceeding in a manner similar to Example 27(m) above but using L-N-boc-benzylsulfonylalanine and (S)-2-amino-1-benzoxazol-2-yl-pentan-1-ol {200mg 0.91mmol, Reference Example 17(c)} there was prepared {(R)-1-[1-(Benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester. LC/MS $m/z=546(M+H)$, 490 (M+H-isobutene)

(q) {(R)-1-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-3-phenyl-propylcarbamoyl]-2-cyclopropylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester

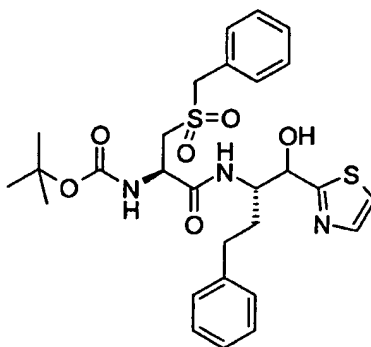


By proceeding in a manner similar to Example 27(m) above but using (2S)-2-amino-4-phenyl-1-benzoxazol-2-yl-butan-1-ol there was prepared {(R)-1-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-3-

-135-

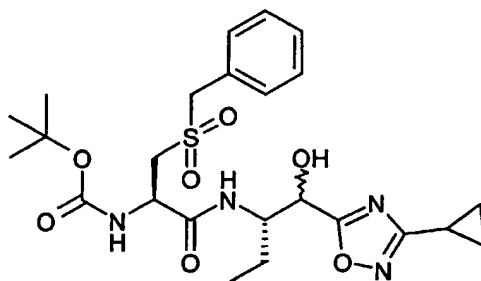
phenyl-propylcarbamoyl]-2-cyclopropylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester.
 LC/MS $m/z=572(M+H)$, 516 ($M+H$ -isobutene).

(r) {(R)-1-[(S)-1-(Hydroxy-thiazol-2-yl-methyl)-3-phenyl-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester



By proceeding in a manner similar to Example 27(m) above but using L-N-boc-benzylsulfonylalanine and (2S)-2-amino-4-phenyl-1-thiazol-2-yl-butan-1-ol (Reference Example 13) there was prepared {(R)-1-[(S)-1-(Hydroxy-thiazol-2-yl-methyl)-3-phenyl-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester. LC/MS $m/z=574(M+H)$

(s) {(R)-2-phenylmethanesulfonyl-1-[(S)-1-[(3-cyclopropyl-1,2,4-oxadiazol-5-yl)-hydroxy-methyl]-propylcarbamoyl]-ethyl}-carbamic acid tert-butyl ester

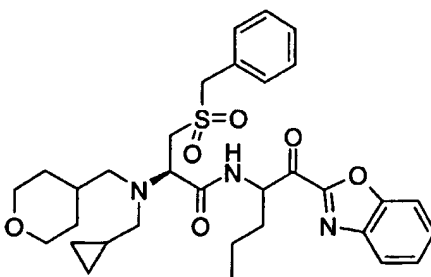


By proceeding in a manner similar to Example 27(m) above but using L-N-boc-benzylsulfonylalanine and (S)-2-amino-1-(3-cyclopropyl-1,2,4-oxadiazol-5-yl)-butan-1-ol (Reference Example 14) there was prepared {(R)-2-phenylmethanesulfonyl-1-[(S)-1-[(3-cyclopropyl-1,2,4-oxadiazol-5-yl)-hydroxy-methyl]-propylcarbamoyl]-ethyl}-carbamic acid tert-butyl ester.

EXAMPLE 28

(R)-N-[1-(Benzoxazole-2-carbonyl)-butyl]-2-[cyclopropylmethyl-(tetrahydro-pyran-4-ylmethyl)-amino]-3-phenylmethanesulfonyl-propionamide

-136-

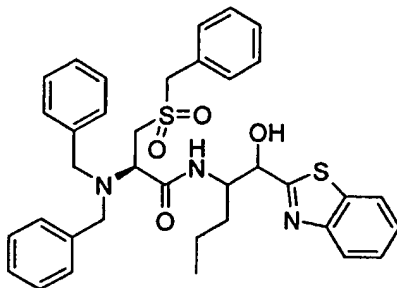


- Step 1. (R)-2-Amino-N-[1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-propionamide {200mg, 0.448mmol, Reference Example 11(i)} was dissolved in 5% acetic acid in acetonitrile (10ml). Tetrahydro-pyran-4-carbaldehyde (51mg, 0.448mmol) was added and the reaction mixture stirred for 16 hours. (Polystyrylmethyl)trimethylammonium cyanoborohydride (218mg, 0.896mmol) was added and the reaction mixture stirred for 3 hours. Cyclopropanecarbaldehyde (157mg, 2.24mmol) was added and stirring continued for 3 hours. The mixture was filtered under suction and the filtrate concentrated under high vacuum.
- Step 2. The residue was dissolved in 10ml dichloromethane. The Dess-Martin-periodinane (380mg, 0.896mmol) was added and the resulting reaction mixture stirred for two hours. The reaction mixture was poured into a 1/1-mixture of saturated sodium bicarbonate solution and saturated sodium thiosulfate solution. The aqueous phase was extracted with dichloromethane. The combined organic phases were washed with saturated sodium bicarbonate solution and brine. The organic phase was dried with magnesium sulfate and the dichloromethane evaporated under reduced pressure. The crude product was purified via flash chromatography (heptane/ethyl acetate 2/1 followed by heptane/ethyl acetate 1/1 to elute) to give (R)-N-[1-(benzoxazole-2-carbonyl)-butyl]-2-(cyclopropylmethyl-(tetrahydro-pyran-4-ylmethyl)-amino]-3-phenylmethanesulfonyl-propionamide as mixture of diastereomers. (83mg, 0.139mmol, 31%). LC/MS $m/z=596$ (M+H) retention time 3.84 (method C).

20

EXAMPLE 29

- (a) (R)-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-2-dibenzylamino-3-phenylmethanesulfonyl-propionamide



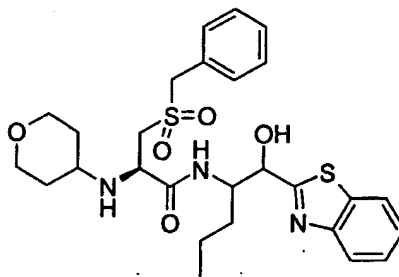
25

(R)-2-Amino-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-

-137-

propionamide {50mg, 0.11mmol, Reference Example 11(a)} was dissolved in a mixture of acetonitrile (5ml) and acetic acid (1ml). Benzaldehyde (56 μ l, 0.55mmol, 5 equivalents) and resin bound cyanoborohydride (54mg, 0.22mmol, 2 equivalents) were added. The reaction mixture was stirred overnight, filtered under suction and the filtrate evaporated to give the (R)-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-2-dibenzylamino-3-phenylmethanesulfonyl-propionamide which was used without further purification in the preparation of Example 18(c).

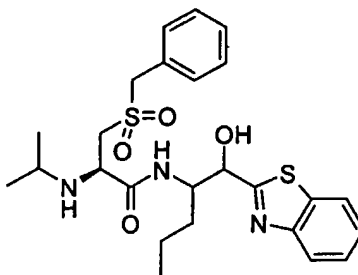
(b) (R)-N-[1-(Benzothiazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide



By proceeding in a manner similar to Example 29(a) above but using tetrahydro-4H-pyran-4-one (51 μ l, 0.55mmol, 5 equivalents) there was prepared (R)-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide. LC/MS m/z =546 (M+H)

EXAMPLE 30

(a) (R)-N-[1-(Benzothiazol-2-yl-hydroxy-methyl)-butyl]-2-isopropylamino-3-phenylmethanesulfonyl-propionamide



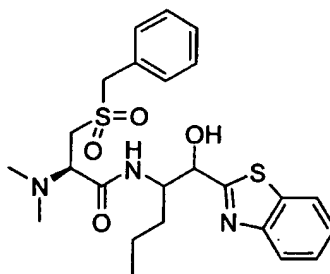
(R)-2-Amino-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-propionamide {50mg, 0.11mmol, Reference Example 11(a)} was dissolved in a mixture of acetonitrile (5ml) and acetic acid (1ml). Acetone (500 μ l) and resin bound cyanoborohydride (54mg, 0.22mmol, 2 equivalents) were added. The reaction mixture was stirred overnight, filtered under suction and

-138-

concentrated under vacuum. The residue was dissolved in dichloromethane and AP Trisamine (Argonaut Technology) (550mg, 1.2mmol) was added. The mixture was stirred for two hours, filtered under suction and the filtrate concentrated under vacuum to give (R)-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-2-isopropylamino-3-phenylmethanesulfonyl-propionamide (30mg, 0.06mmol, 54%).

5 LC/MS $m/z=504$ (M+H).

(b) (R)-N-[1-(Benzothiazol-2-yl-hydroxy-methyl)-butyl]-2-dimethylamino-3-phenylmethanesulfonyl-propionamide

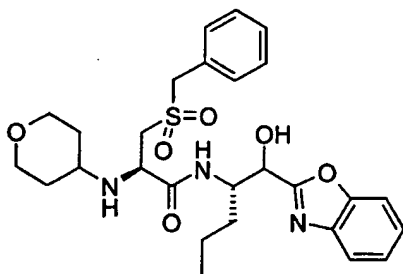


10 By proceeding in a manner similar to Example 30(a) above but using formaldehyde solution (75 μ l, 1mmol, 37w-% aqueous solution) there was prepared (R)-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-2-dimethylamino-3-phenylmethanesulfonyl-propionamide (30mg, 54%). LC/MS $m/z=490$ (M+H).

15

EXAMPLE 31

(a) (R)-N-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide



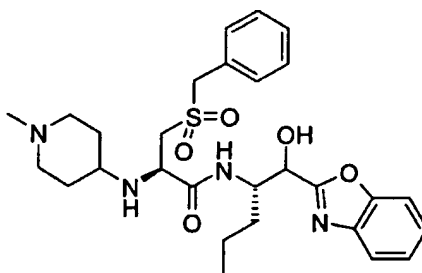
A solution of (R)-2-amino-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-propionamide {100mg, 0.22mmol, Reference Example 11(c)} in a mixture of acetonitrile (5mL) and acetic acid (1mL) was treated with tetrahydro-4H-pyran-4-one (101 μ l, 1.1mmol). After agitating at room temperature for 3 hours the mixture was then treated with resin-bound cyanoborohydride (108mg, 0.44mmol) and agitation was continued overnight. The reaction mixture was filtered and the filtrate was evaporated. The residue was dissolved in dichloromethane (10mL) and the solution was treated with Silicycle Triamine (611mg, 2.2mmol), then agitated for 2

25

-139-

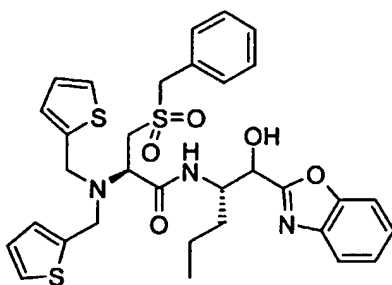
hours and then filtered. The solution of (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide was used directly in the preparation of Example 20(b).

- 5 (b) (R)-N-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-(1-methyl-piperidin-4-ylamino)-3-phenylmethanesulfonyl-propionamide



- 10 By proceeding in a manner similar to Example 31(a) above but using 1-methyl-4-piperidone (136μl, 1.1mmol) there was prepared (R)-N-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-(1-methyl-piperidin-4-ylamino)-3-phenylmethanesulfonyl-propionamide was used directly in the preparation of Example 19(b).

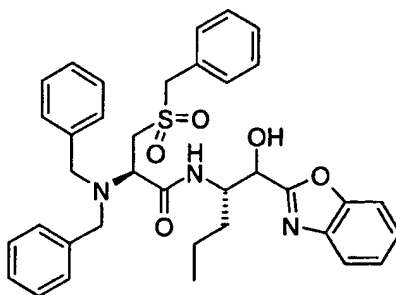
- 15 (c) (R)-N-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-(bis-thiophen-2-ylmethyl-amino)-3-phenylmethanesulfonyl-propionamide



- 20 By proceeding in a manner similar to Example 31(a) above but using 2-thiophenecarboxaldehyde (20μl, 0.22mmol) there was prepared (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-(bis-thiophen-2-ylmethyl-amino)-3-phenylmethanesulfonyl-propionamide was used directly in the preparation of Example 19(c).

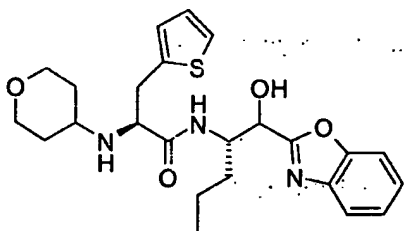
- (d) (R)-N-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-dibenzylamino-3-phenylmethanesulfonyl-propionamide

-140-



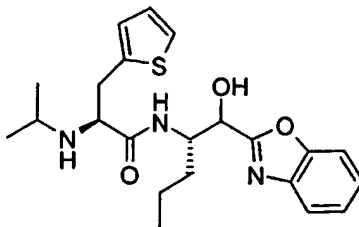
By proceeding in a manner similar to Example 31(a) above but using benzaldehyde (22 μ l, 0.22mmol) there was prepared (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-dibenzylamino-3-phenylmethanesulfonyl-propionamide which was used directly in the preparation of Example 19(d).

(e) (S)-N-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-(tetrahydro-pyran-4-ylamino)-3-thiophen-2-yl-propionamide



By proceeding in a manner similar to Example 317(a) above but using (S)-2-amino-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-thiophen-2-yl-propionamide {82mg, 0.22mmol, Reference Example 11(b)} and tetrahydro-4H-pyran-4-one (101 μ l, 1.1mmol) there was prepared (S)-N-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-(tetrahydro-pyran-4-ylamino)-3-thiophen-2-yl-propionamide which was used directly in the preparation of Example 19(e).

(f) (S)-N-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-isopropylamino-3-thiophen-2-yl-propionamide

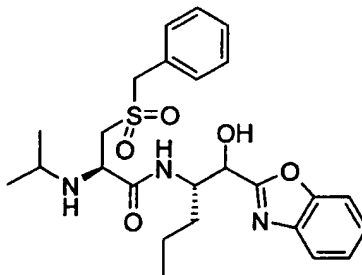


By proceeding in a manner similar to Example 31(a) above but using (S)-2-amino-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-thiophen-2-yl-propionamide {82mg, 0.22mmol, Reference

-141-

Example 11(b)} and acetone (100 μ l) there was prepared (S)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-isopropylamino-3-thiophen-2-yl-propionamide which was used directly in the preparation of Example 19(f).

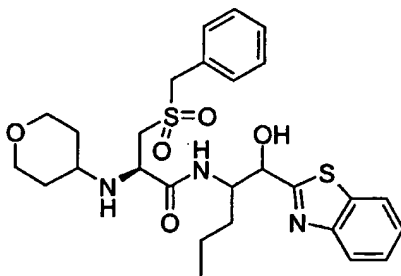
- 5 (g) (R)-N-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-isopropylamino-3-phenylmethanesulfonyl-propionamide



- 10 By proceeding in a manner similar to Example 31(a) above but using acetone (500 μ l) there was prepared (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-isopropylamino-3-phenylmethanesulfonyl-propionamide (30.5mg, 29%). LC/MS m/z =488 (M+H).

EXAMPLE 32

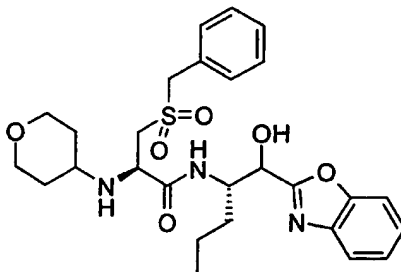
- (a) (R)-N-[1-(Benzothiazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide



- 15 A solution of (R)-2-amino-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-propionamide {100mg, 0.22mmol, Reference Example 11(a)} in a mixture of acetonitrile and acetic acid (10mL, 95:5, v/v) was treated with tetrahydro-4H-pyran-4-one (101 μ l, 1.1mmol) and resin-bound cyanoborohydride (108mg, 0.44mmol). This mixture was stirred at room temperature overnight then
20 evaporated. The residue was dissolved in dichloromethane and the solution was treated with Silicycle Triamine (611mg, 2.2mmol) then stirred at room temperature for 2 hours then filtered. The filtrate was evaporated to give (R)-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide, LC/MS m/z =546 (M+H), which was used directly in the preparation of Example 18(b).

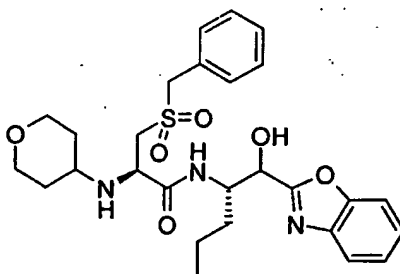
-142-

- (b) (R)-N-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide



By proceeding in a manner similar to Example 32(a) above but using (R)-2-amino-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-propionamide {98mg, 0.22mmol, Reference Example 11(c)} there was prepared (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide, LC/MS $m/z=530$ (M+H), which was used directly in the preparation of Example 19(a).

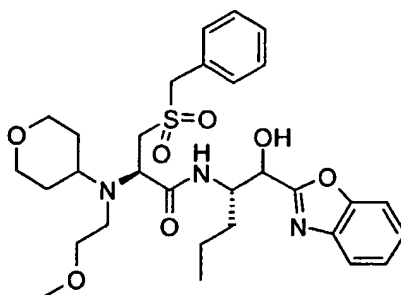
- 10 (c) (R)-N-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide



- By proceeding in a manner similar to Example 32(a) above but using (R)-2-amino-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-propionamide {Reference Example 11(c)} there was prepared (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide (106mg, 91%). LC/MS $m/z=530$ (M+H).

- 20 (d) (R)-N-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-[(2-methoxy-ethyl)-(tetrahydro-pyran-4-yl)-amino]-3-phenylmethanesulfonyl-propionamide

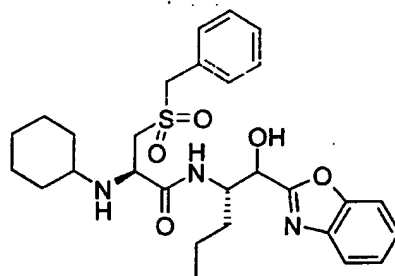
-143-



By proceeding in a manner similar to Example 32(a) above but using (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide {53mg, 0.1mmol, Reference Example 32(c)} and 2-methoxyethanal (53mg, 0.55mmol) there was prepared (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-[(2-methoxy-ethyl)-(tetrahydro-pyran-4-yl)-amino]-3-phenylmethanesulfonyl-propionamide (56mg, 95%).

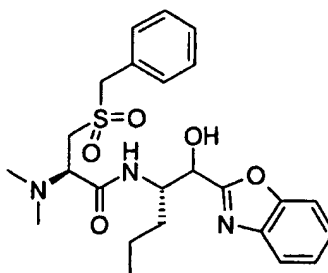
LC/MS m/z =588 (M+H)

(e) (R)-N-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-cyclohexylamino-3-phenylmethanesulfonyl-propionamide



By proceeding in a manner similar to Example 32(a) above but using (R)-2-amino-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-propionamide {49mg, 0.11mmol, Reference 11(c)} and cyclohexanone (52μl, 0.5mmol) there was prepared (R)-N-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-cyclohexylamino-3-phenylmethanesulfonyl-propionamide (48mg, 83%).

(f) (R)-N-[(S)-1-(Benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-dimethylamino-3-phenylmethanesulfonyl-propionamide



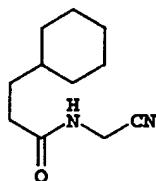
-144-

By proceeding in a manner similar to Example 32(a) above but using (R)-2-amino-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-propionamide {49mg, 0.11mmol, Reference Exaple 11(c)} and formaldehyde (75 μ l, 1mmol, 37 w-% in water), there was prepared (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-dimethylamino-3-phenylmethanesulfonyl-propionamide (10mg, 19%). LC/MS m/z=474 (M+H).

EXAMPLE 33

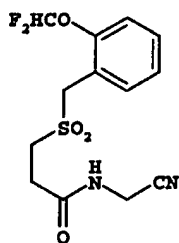
The following compounds of Formula 1 are provided by methods described in the application:

10 (a) N-Cyanomethyl-3-cyclohexyl-propionamide



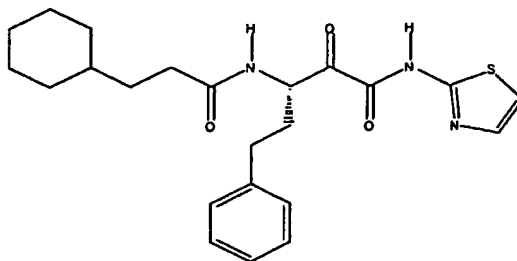
¹H NMR: (CDCl₃) 6.22 (br s, 1H), 4.20 (s, 2H), 2.23 (m, 2H), 1.65 (m, 5H), 1.50 (m, 2H), 1.10-1.30 (m, 4H), 0.90 (m, 2H); LC-MS: t=3.67min., 193.0(M-1), 195.1(M+1). MS: API 150EX. (LC: Agilent 1100Series, Column: Phenomenex, 5u ODS3 100A 100X3mm. Flow Rate: 2ml/min. Two solvent gradient: Solvent A, 99% water, 1% acetonitrile, 0.1% AcOH. Solvent B, 99% acetonitrile, 1% water, 0.1% AcOH. Gradient from 100% A, 0% B to 0% A, 100% B from t = 0 to t = 6min. Then gradient back to 100% A, 0% B from t = 7 to t = 15 min.);

15 (b) N-Cyanomethyl-3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionamide

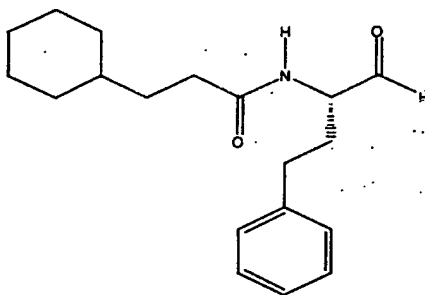


20 ¹H NMR: (CDCl₃) 7.52 (d, 1H, J=8Hz), 7.43 (t, 1H, J=8Hz), 7.29 (d, 1H, J=8Hz), 7.20 (d, 1H, J=8Hz), 6.40 (m, 1H), 4.41 (s, 2H), 4.16 (d, 2H, J=6Hz), 3.72 (s, 1H), 3.34 (t, 2H, J=8Hz), 2.77 (t, 2H, J=8Hz); LC-MS: t=3.02min., 331.1(M-1), 333.1(M+1). MS: API 150EX. (LC: Agilent 1100Series, Column: Phenomenex, 5u ODS3 100A 100X3mm. Flow Rate: 2ml/min. Two solvent gradient: Solvent A, 99% water, 1% acetonitrile, 0.1% AcOH. Solvent B, 99% acetonitrile, 1% water, 0.1% AcOH. Gradient from 100% A, 0% B to 0% A, 100% B from t = 0 to t = 6min. Then gradient back to 100% A, 0% B from t = 7 to t = 15 min.).

-145-

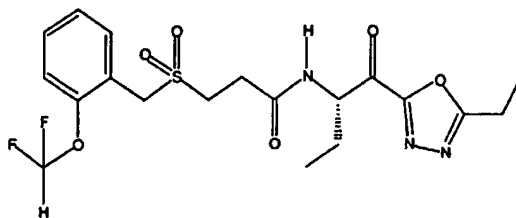
(c) 3-(3-Cyclohexyl-propionylamino)-2-oxo-5-phenyl-pentanoic acid thiazol-2-ylamide

data for the compound as drawn and for it's enol and hydrate forms: LC-MS: $t_r=4.74$ min. 426.4(M-1), 428.2(M+1); 4.97min, 426.2 (M-1), 428.2 (M+1); 5.57min, 426.3(M-1), 427.9 (M+1). MS: API 150EX. (LC: Agilent 1100Series, Column: Phenomenex, 5u ODS3 100A 100X3mm. Flow Rate: 2ml/min. Two solvent gradient: Solvent A, 99% water, 1% acetonitrile, 0.1% AcOH. Solvent B, 99% acetonitrile, 1% water, 0.1% AcOH. Gradient from 100% A, 0% B to 0% A, 100% B from $t = 0$ to $t = 6$ min. Then gradient back to 100% A, 0% B from $t = 7$ to $t = 15$ min.)

10 (d) 3-Cyclohexyl-N-(1-formyl-3-phenyl-propyl)-propionamide

LC-MS: $t_r=4.57$ min., 300.4(M-1), 302.3(M+1). MS: API 150EX. (LC: Agilent 1100Series, Column: Phenomenex, 5u ODS3 100A 100X3mm. Flow Rate: 2ml/min. Two solvent gradient: Solvent A, 99% water, 1% acetonitrile, 0.1% AcOH. Solvent B, 99% acetonitrile, 1% water, 0.1% AcOH. Gradient from 100% A, 0% B to 0% A, 100% B from $t = 0$ to $t = 6$ min. Then gradient back to 100% A, 0% B from $t = 7$ to $t = 15$ min.)

(f) 3-(2-Difluoromethoxy-phenylmethanesulfonyl)-N-[(S)-1-(5-ethyl-[1,3,4]oxadiazole-2-carbonyl)-propyl]-propionamide



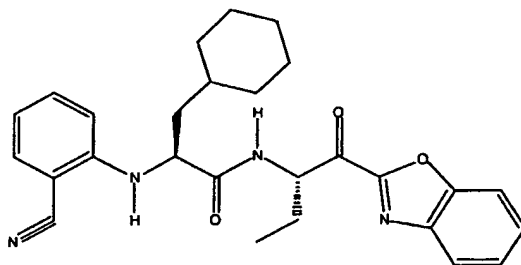
LC-MS: $R_T=2.32$ min., 460.3(M+1) 482.2(M+23) MS: API 150EX. (LC: Agilent 1100Series, Column: Phenomenex, 5u ODS3 100A 100X3mm. Flow Rate: 2ml/min. Two solvent gradient:

-146-

Solvent A, 99% water, 1% acetonitrile, 0.1% AcOH. Solvent B, 99% acetonitrile, 1% water, 0.1% AcOH. Gradient from 100% A, 0% B to 0% A, 100% B from $t = 0$ to $t = 2.5$ min. Then gradient back to 100% A, 0% B from $t = 3.0$ to $t = 3.5$ min. Then gradient held at 100% A, 0% B from $t = 3.5$ to 5 min.)

5

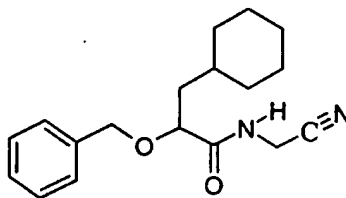
(g) *N*-[(*S*)-1-(Benzooxazole-2-carbonyl)-propyl]-2-(2-cyano-phenylamino)-3-cyclohexyl-propionamide



^1H NMR: (CDCl_3) 7.83 (d, 1H, $J=8\text{Hz}$), 7.59 (d, 1H, $J=8\text{Hz}$), 7.43-7.58 (m, 2H), 7.02-7.25 (m, 4H), 6.59 (t, 1H, $J=8\text{Hz}$), 6.49 (d, 1H, $J=8\text{Hz}$), 5.40-5.47 (m, 1H), 4.77 (m, 1H), 3.83-3.88 (m, 1H), 2.12-2.22 (m, 1H), 1.85-2.00 (m, 2H), 1.55-1.83 (m, 8H), 1.12-1.35 (m, 4H), 0.95-1.10 (m, 3H); LC-MS: $t=2.97\text{min.}$, 457.5($M-1$), 459.3($M+1$), 481.4($M+23$) MS: API 150EX. (LC: Agilent 1100Series, Column: Phenomenex, 5u ODS3 100A 100X3mm. Flow Rate: 2ml/min. Two solvent gradient: Solvent A, 99% water, 1% acetonitrile, 0.1% AcOH. Solvent B, 99% acetonitrile, 1% water, 0.1% AcOH. Gradient from 100% A, 0% B to 0% A, 100% B from $t = 0$ to $t = 2.5$ min. Then gradient back to 100% A, 0% B from $t = 3.0$ to $t = 3.5$ min. Then gradient held at 100% A, 0% B from $t = 3.5$ to 5 min.)

(h) *N*-Cyanomethyl-3-cyclohexyl-2-(4-methoxy-phenoxy)-propionamide (Compound 1); ^1H NMR: (CDCl_3) 7.42-7.36 (m, 5H), 6.90 (t, 1H), 4.55 (d, 1H), 4.51 (d, 1H), 4.22 (dd, 1H), 4.16 (dd, 1H), 4.00 (t, 1H), 1.70-0.80 (m, 13H); MS: (M^++1) 301;

(i) 2-Benzoyloxy-*N*-cyanomethyl-3-cyclohexyl-propionamide (Compound 2)



25 using 2(R)-benzyloxy-4-phenyl-butyric acid as starting material. ^1H NMR: (CDCl_3) δ 6.84-6.80 (m, 4H), 6.75 (t, 1H), 4.55 (dd, 1H), 4.24 (dd, 1H), 4.12 (dd, 1H), 3.78 (s, 3H), 1.80-0.85 (m, 13H); MS: ($M-1$) 315.

-147-

- (j) (R)-N-[(S)-1-(1-benzooxazol-2-yl-methanoyl)-butyl]-2-benzyloxy-3-phenylmethanesulfonyl-propionamide (Compound 3); ¹H NMR: (CDCl₃) 7.89 (d, 1H), 7.68 (d, 1H), 7.60-7.32 (m, 13H), 5.70 (m, 1H), 4.79 (d, 1H), 4.77 (d, 1H), 4.53 (dd, 1H), 4.33 (d, 1H), 4.30 (d, 1H), 3.38 (dd, 1H), 3.25 (dd, 1H), 2.15-2.05 (m, 1H), 1.84-75 (m, 1H), 1.45-1.30 (m, 2H), 0.93 (t, 3H); MS: (M⁺+1) 535, (M-1) 533;
- (k) (R)-N-[(S)-1-(1-benzooxazol-2-yl-methanoyl)-propyl]-2-methoxymethoxy-3-phenylmethanesulfonyl-propionamide (Compound 9); ¹HNMR (DMSO): 8.87(d, J=6.91Hz, 1H), 7.99(d, J=7.91Hz, 1H), 7.89(d, J=8.15Hz, 1H), 7.64(t, J=8.1Hz, 1H), 7.54(t, J=8.1Hz, 1H), 7.4-7.3(m, 5H), 5.3-5.2(m, 1H), 4.7-4.65(m, 1H), 4.65-4.63(m, 2H), 4.55-4.50(m, 2H), 3.53-3.26(m, 2H), 3.34(s, 3H), 2.11-1.98(m, 1H), 1.81-1.69(m, 1H), 0.97(t, J=7.15Hz, 3H); MS: 473(M-1), 497(M+23);
- (l) (S)-N-[(S)-1-(1-benzooxazol-2-yl-methanoyl)-butyl]-2-hydroxy-3-phenyl-propionamide (Compound 10);
- (m) (R)-N-[(S)-1-(1-benzooxazol-2-yl-methanoyl)-propyl]-3-phenylmethanesulfonyl-2-triisopropylsilyloxy-propionamide (Compound 12); ¹HNMR (CD₃Cl): 7.93(d, J=8.15Hz, 1H), 7.6(d, J=8.1Hz, 1H), 7.6-7.4(m, 3H), 7.4-7.3(m, 5H), 5.85-5.73(m, 1H), 4.85-4.74(m, 1H), 4.5-4.3(m, 2H); 3.47-3.35(m, 2H), 2.35-2.15(m, 1H), 2.15-1.95(m, 1H), 1.3-0.8(m, 24H); MS: 609.4(M+23);
- (n) (R)-N-[(S)-1-(1-benzothiazol-2-yl-methanoyl)-propyl]-2-hydroxy-3-phenylmethanesulfonyl-propionamide (Compound 13); ¹HNMR (CD₃Cl): 8.21(d, J=8.67Hz, 1H), 7.98(d, J=8.6Hz, 1H), 7.7-7.55(m, 3H), 7.45-7.3(m, 5H), 5.8-5.7(m, 1H), 4.75-4.6(m, 1H), 4.4-4.3(m, 2H), 4.08(br, 1H), 3.62-3.5(m, 1H), 3.3-3.1(m, 1H), 2.3-2.15(m, 1H), 2.05-1.9(m, 1H), 0.997(t, J=7.4Hz, 3H); MS: 469.2(M+23);
- (o) (R)-2-hydroxy-3-phenylmethanesulfonyl-N-[(S)-1-(1-pyridazin-3-yl-methanoyl)-butyl]-propionamide (Compound 16); ¹HNMR (CD₃Cl): 9.35(dd, J=4.93Hz, J=1.72Hz, 1H), 8.14(dd, J=1.72Hz, J=8.39Hz, 1H), 7.69(dd, J=4.93Hz, J=8.39Hz, 1H), 7.65(d, J=7.6Hz, 1H), 7.5-7.36(m, 5H), 6.04-5.96(m, 1H), 4.75-4.63(m, 1H), 4.45-4.3(m, 3H), 3.53(dd, J=2.48Hz, J=14.85Hz, 1H), 3.22(dd, J=14.82Hz, J=2.48Hz, 1H), 2.2-2.07(m, 1H), 1.81-1.65(m, 1H), 1.6-1.2(m, 2H), 0.93(t, J=7.18Hz, 3H); MS: 403.6(M-1), 428(M+23);
- (p) (S)-3-((R)-2-hydroxy-3-phenylmethanesulfonyl-propanoylamino)-2-oxo-pentanoic acid benzylamide (Compound 18); ¹HNMR (CD₃Cl): 7.45-7.25(m, 10H), 5.34-5.26(m, 1H), 4.7-4.6(m,

-148-

1H), 4.47(d, J=6.18Hz, 2H), 4.4-4.3(m, 2H), 4.15-4.05(m, 1H), 3.55-3.45(m, 1H), 3.25-3.13(m, 1H), 2.2-2.0(m, 1H), 1.8-1.6(m, 1H), 1.61(s, 2H), 0.95(t, J=6.91Hz, 3H); MS: 469.2(M+23);

- (q) (R)-N-[(S)-1-(1-benzooxazol-2-yl-methanoyl)-propyl]-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionamide (Compound 21); ¹HNMR (CD₃Cl): 7.91(d, J=7.91Hz, 1H), 7.75(d, J=7.9Hz, 1H), 7.7-7.2(m, 6H), 6.63(t, J=73.41Hz, 1H), 5.7-5.58(m, 1H), 5.4-5.29(m, 1H), 4.7-4.6(m, 1H), 4.51(s, 2H), 4.19(br, 1H), 3.72-3.63(m, 1H), 3.35-3.2(m, 1H), 2.3-2.0(m, 1H), 2.0-1.7(m, 1H), 0.99(t, J=6.9Hz, 3H); MS: 495.5(M-1), 497.2(M+1);
- (r) (R)-N-[(S)-1-(1-benzothiazol-2-yl-methanoyl)-propyl]-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionamide (Compound 22); ¹HNMR (CD₃Cl): 8.21(d, J=8.15Hz, 1H), 7.99(d, J=8.1Hz, 1H), 7.73-7.2(m, 6H), 6.63(t, J=73.4Hz, 1H), 5.85-5.75(m, 1H), 5.3(s, 1H), 4.78-4.7(m, 1H), 4.56-4.4(m, 2H), 4.19-4.09(m, 1H), 3.7-3.6(m, 1H), 3.35-3.2(m, 1H), 2.28(s, 2H), 1.27(t, J=6.9Hz, 3H); MS: 511.4(M-1), 513.6(M+1); and
- (s) (2R,5S)-2-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonylmethyl]-6-ethoxy-5-ethyl-morpholin-3-one (Compound 24).

ENZYME ASSAY EXAMPLE

Cathepsin S Assay

- Solutions of test compounds in varying concentrations were prepared in 10 μ L of dimethyl sulfoxide (DMSO) and then diluted into assay buffer (40 μ L, comprising: MES, 50 mM (pH 6.5); EDTA, 2.5 mM; and NaCl, 100 mM). Human cathepsin S (0.158 pMoles in 25 μ L of assay buffer) was added to the dilutions. The assay solutions were mixed for 5-10 seconds on a shaker plate, covered and incubated for 30 minutes at ambient temperature. Z-Val-Val-Arg-AMC (9 nMoles in 25 μ L of assay buffer) was added to the assay solutions and hydrolysis was followed spectrophotometrically at (λ 460 nm) for 5 minutes. Apparent inhibition constants (K_i) were calculated from the enzyme progress curves using standard mathematical models.

ENZYME ASSAY EXAMPLE

Cathepsin B Assay

- Solutions of test compounds in varying concentrations were prepared in 10 μ L of dimethyl sulfoxide (DMSO) and then diluted into assay buffer (40 μ L, comprising: *N,N*-bis(2-hydroxyethyl)-2-aminoethanesulfonic acid (BES), 50 mM (pH 6); polyoxyethylenesorbitan monolaurate, 0.05%; and dithiothreitol (DTT), 2.5 mM). Human cathepsin B (0.025 pMoles in 25 μ L of assay buffer) was added to the dilutions. The assay solutions were mixed for 5-10 seconds on a shaker plate, covered and incubated for 30 minutes at ambient temperature. Z-FR-AMC (20 nMoles in 25 μ L of assay buffer) was added to the assay solutions and hydrolysis was followed

-149-

spectrophotometrically at (λ 460 nm) for 5 minutes. Apparent inhibition constants (K_i) were calculated from the enzyme progress curves using standard mathematical models.

5

ENZYME ASSAY EXAMPLE

Cathepsin K Assay

Solutions of test compounds in varying concentrations were prepared in 10 μ L of dimethyl sulfoxide (DMSO) and then diluted into assay buffer (40 μ L, comprising: MES, 50 mM (pH 5.5); EDTA, 2.5 mM; and
10 DTT, 2.5 mM). Human cathepsin K (0.0906 pMoles in 25 μ L of assay buffer) was added to the dilutions. The assay solutions were mixed for 5-10 seconds on a shaker plate, covered and incubated for 30 minutes at ambient temperature. Z-Phe-Arg-AMC (4 nMoles in 25 μ L of assay buffer) was added to the assay solutions and hydrolysis was followed spectrophotometrically at (λ 460 nm) for 5 minutes. Apparent inhibition constants (K_i) were calculated from the enzyme progress curves using standard mathematical models.

15

ENZYME ASSAY EXAMPLE

Cathepsin L Assay

Solutions of test compounds in varying concentrations were prepared in 10 μ L of dimethyl sulfoxide (DMSO) and then diluted into assay buffer (40 μ L, comprising: MES, 50 mM (pH 5.5); EDTA, 2.5 mM; and
20 DTT, 2.5 mM). Human cathepsin L (0.05 pMoles in 25 μ L of assay buffer) was added to the dilutions. The assay solutions were mixed for 5-10 seconds on a shaker plate, covered and incubated for 30 minutes at ambient temperature. Z-Phe-Arg-AMC (1 nMoles in 25 μ L of assay buffer) was added to the assay solutions and
25 hydrolysis was followed spectrophotometrically at (λ 460 nm) for 5 minutes. Apparent inhibition constants (K_i) were calculated from the enzyme progress curves using standard mathematical models.

According to applicants' assays conducted as described above, the apparent inhibition constants (K_i) for the following listed compounds of the invention, against Cathepsin S, were about or below 0.01 μ M:

30 morpholine-4-carboxylic acid (*R*)-1-(cyanomethyl-carbamoyl)-2-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-ethyl ester, (Compound 31), Example 3(a);

morpholine-4-carboxylic acid (*R*)-1-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester, (Compound 11), Example 4(a);

35

morpholine-4-carboxylic acid (*R*)-1-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propylcarbamoyl]-2-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-ethyl ester, (Compound 14), Example 4(b);

40

morpholine-4-carboxylic acid (*R*)-1-[(*S*)-1-(1-benzothiazol-2-yl-methanoyl)-propylcarbamoyl]-2-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-ethyl ester, (Compound 15), Example 4(c);

pyrrolidine-1-carboxylic acid (*R*)-1-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester, (Compound 19). Example 4(d);

-150-

dimethyl-carbamic acid (*R*)-1-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester, (Compound 20), Example 4(e);

5 morpholine-4-carboxylic acid (*R*)-1-[(*S*)-1-(1-benzylcarbamoyl-methanoyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester, (Compound 25), Example 4(f);

morpholine-4-carboxylic acid (*S*)-1-[(*S*)-1-(oxazolo[4,5-*b*]pyridine-2-carbonyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester, Example 4(g);

10 morpholine-4-carboxylic acid (*S*)-1-[(*S*)-1-(5-ethyl-[1,3,4]oxadiazole-2-carbonyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester, Example 4(h);

(*R*)-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-*N*-[(*S*)-1-formyl-propyl]-2-hydroxy-propionamide, (Compound 23), Example 6;

15 (*R*)-*N*-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propyl]-2-hydroxy-3-phenyl-methanesulfonyl-propionamide, (Compound 5), Example 7;

20 (*S*)-3-{3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-propanoylamino}-2-oxo-pentanoic acid benzylamide, (Compound 27), Example 8(a);

(*R*)-*N*-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propyl]-2-(2-nitro-phenylamino)-3-phenylmethanesulfonyl-propionamide, (Compound 28), Example 9;

25 (*R*)-*N*-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-butyl]-2-(5-nitro-thiazol-2-ylamino)-3-phenylmethanesulfonyl-propionamide, (Compound 29), Example 10;

(*R*)-*N*-[(*S*)-1-(benzoxazole-2-carbonyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide; Example 19(a);

30 (*R*)-*N*-[(*S*)-1-(benzoxazole-2-carbonyl)-butyl]-2-isopropylamino-3-phenylmethanesulfonyl propionamide, Example 21(a);

35 (*R*)-*N*-[(*S*)-1-(benzoxazole-2-carbonyl)-butyl]-2-[(2-methoxy-ethyl)-(tetrahydro-pyran-4-yl)-amino]-3-phenylmethanesulfonyl-propionamide, Example 21(b);

(*R*)-*N*-[(*S*)-1-(benzoxazole-2-carbonyl)-butyl]-2-cyclohexylamino-3-phenylmethanesulfonyl-propionamide, Example 21(c);

40 morpholine-4-carboxylic acid (*S*)-2-cyclohexyl-1-[(*S*)-1-(oxazolo[4,5-*b*]pyridine-2-carbonyl)-propylcarbamoyl]-ethyl ester, Example 24(b);

3-(2-difluoromethoxy-phenylmethanesulfonyl)-*N*-[(*S*)-1-(oxazolo[4,5-*b*]pyridine-2-carbonyl)-propyl]-propionamide, Example 33(e);

45 (*S*)-3-((*R*)-2-hydroxy-3-phenylmethanesulfonyl-propanoylamino)-2-oxo-pentanoic acid benzylamide (Compound 18), Example 33(p);

50 (*R*)-*N*-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propyl]-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionamide (Compound 21), Example 33(q);

Moreover, the compounds of the present invention were observed to have varying degrees of selective inhibitory action on cathepsin S protease. For example, the above listed 22 compounds were found to inhibit

-151-

cathepsin S protease activity at concentrations that are more than 75 fold less than those concentrations required to produce an equiactive inhibition on cathepsin K protease.

-152-

EXAMPLE

Representative Pharmaceutical Formulations Containing a Compound of Formula I

ORAL FORMULATION

5	Compound of Formula I	10-100 mg
	Citric Acid Monohydrate	105 mg
	Sodium Hydroxide	18 mg
	Flavoring	
	Water	q.s. to 100 mL

10

INTRAVENOUS FORMULATION

	Compound of Formula I	0.1-10 mg
	Dextrose Monohydrate	q.s. to make isotonic
	Citric Acid Monohydrate	1.05 mg
15	Sodium Hydroxide	0.18 mg
	Water for Injection	q.s. to 1.0 mL

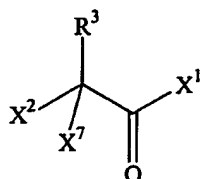
TABLET FORMULATION

	Compound of Formula I	1%
20	Microcrystalline Cellulose	73%
	Stearic Acid	25%
	Colloidal Silica	1%

-153-

WE CLAIM:

1. A compound of Formula I:



I

in which:

 X^1 is $-NHC(R^1)(R^2)X^3$ or $-NHX^4$; X^2 is hydrogen, fluoro, $-OH$, $-OR^4$, $-NHR^{15}$ or $-NR^{17}R^{18}$ and X^7 is hydrogen or X^2 and10 X^7 both represent fluoro; X^3 is cyano, $-C(R^7)(R^8)R^{16}$, $-C(R^6)(OR^6)_2$, $-CH_2C(O)R^{16}$, $-CH=CHS(O)_2R^5$, $-C(O)CF_2C(O)NR^5R^5$, $-C(O)C(O)NR^5R^6$, $-C(O)C(O)OR^5$, $-C(O)CH_2OR^5$, $-C(O)CH_2N(R^6)SO_2R^5$ or $-C(O)C(O)R^5$; wherein R^5 is hydrogen, (C_{1-4}) alkyl, (C_{3-10}) cycloalkyl (C_{0-6}) alkyl, hetero (C_{3-10}) cycloalkyl (C_{0-3}) alkyl, (C_{6-10}) aryl (C_{0-6}) alkyl,15 hetero (C_{5-10}) aryl (C_{0-6}) alkyl, (C_{9-10}) bicycloaryl (C_{0-6}) alkyl orhetero (C_{8-10}) bicycloaryl (C_{0-6}) alkyl; R^6 is hydrogen, hydroxy or (C_{1-6}) alkyl; or where X^3 contains an $-NR^5R^6$ group, R^5 and R^6 together with the nitrogen atom to which they are bothattached, form hetero (C_{3-10}) cycloalkyl, hetero (C_{5-10}) aryl or hetero (C_{8-10}) bicycloaryl; R^7 ishydrogen or (C_{1-4}) alkyl and R^8 is hydroxy or R^7 and R^8 together form oxo; R^{16} is hydrogen, -20 X^4 , $-CF_3$, $-CF_2CF_2R^9$ or $-N(R^6)OR^6$; R^9 is hydrogen, halo, (C_{1-4}) alkyl, (C_{5-10}) aryl (C_{0-6}) alkyl or (C_{5-10}) heteroaryl (C_{0-6}) alkyl, with the proviso that when X^3 is cyano, then X^2 is hydrogen,fluoro, $-OH$, $-OR^4$ or $-NR^{17}R^{18}$ and X^7 is hydrogen or X^2 and X^7 both represent fluoro; X^4 comprises a heteromonocyclic ring containing 4 to 7 ring member atoms or a fused heterobicyclic ring system containing 8 to 14 ring member atoms and any carbocyclic ketone,25 iminoketone or thioketone derivative thereof, with the proviso that when $-X^4$ is other than a

heteromonocyclic ring containing 5 ring member atoms, wherein no more than two of the ring

member atoms comprising the ring are heteroatoms, then X^2 is fluoro, $-OH$, $-OR^4$, $-NHR^{15}$ or $-NR^{17}R^{18}$ and X^7 is hydrogen or X^2 and X^7 both represent fluoro;wherein within R^5 , X^3 or X^4 any alicyclic or aromatic ring system is unsubstituted or

substituted further by 1 to 5 radicals independently selected from (C₁₋₆)alkyl, (C₁₋₆)alkylidene, cyano, halo, halo-substituted(C₁₋₄)alkyl, nitro, -X⁵NR¹²R¹², -X⁵NR¹²C(O)R¹², -X⁵NR¹²C(O)OR¹², -X⁵NR¹²C(O)NR¹²R¹², -X⁵NR¹²C(NR¹²)NR¹²R¹², -X⁵OR¹², -X⁵SR¹², -X⁵C(O)OR¹², -X⁵C(O)R¹², -X⁵OC(O)R¹², -X⁵C(O)NR¹²R¹², -X⁵S(O)₂NR¹²R¹², -X⁵NR¹²S(O)₂R¹², -X⁵P(O)(OR¹²)OR¹², -X⁵OP(O)(OR¹²)OR¹², -X⁵NR¹²C(O)R¹³, -X⁵S(O)R¹³ and -X⁵S(O)₂R¹³ and/or 1 radical selected from -R¹⁴, -X⁵OR¹⁴, -X⁵SR¹⁴, -X⁵S(O)R¹⁴, -X⁵S(O)₂R¹⁴, -X⁵C(O)R¹⁴, -X⁵C(O)OR¹⁴, -X⁵OC(O)R¹⁴, -X⁵NR¹⁴R¹², -X⁵NR¹²C(O)R¹⁴, -X⁵NR¹²C(O)OR¹⁴, -X⁵C(O)NR¹²R¹², -X⁵S(O)₂NR¹⁴R¹², -X⁵NR¹²S(O)₂R¹⁴, -X⁵NR¹²C(O)NR¹⁴R¹² and -X⁵NR¹²C(NR¹²)NR¹⁴R¹², wherein X⁵ is a bond or (C₁₋₆)alkylene; R¹² at each occurrence independently is hydrogen, (C₁₋₆)alkyl or halo-substituted(C₁₋₆)alkyl; R¹³ is (C₁₋₆)alkyl or halo-substituted(C₁₋₆)alkyl; and R¹⁴ is (C₃₋₁₀)cycloalkyl(C₀₋₆)alkyl, hetero(C₃₋₁₀)cycloalkyl(C₀₋₃)alkyl, (C₆₋₁₀)aryl(C₀₋₆)alkyl, hetero(C₅₋₁₀)aryl(C₀₋₆)alkyl, (C₉₋₁₀)bicycloaryl(C₀₋₆)alkyl or hetero(C₈₋₁₀)bicycloaryl(C₀₋₆)alkyl;

R¹ is hydrogen or (C₁₋₆)alkyl and R² is selected from a group consisting of hydrogen, cyano, -X⁵NR¹²R¹², -X⁵NR¹²C(O)R¹², -X⁵NR¹²C(O)OR¹², -R¹², -X⁵NR¹²C(O)NR¹²R¹², -X⁵NR¹²C(NR¹²)NR¹²R¹², -X⁵OR¹², -X⁵SR¹², -X⁵C(O)OR¹², -X⁵C(O)R¹², -X⁵OC(O)R¹², -X⁵C(O)NR¹²R¹², -X⁵S(O)₂NR¹²R¹², -X⁵NR¹²S(O)₂R¹², -X⁵P(O)(OR¹²)OR¹², -X⁵OP(O)(OR¹²)OR¹², -X⁵NR¹²C(O)R¹³, -X⁵S(O)R¹³, -X⁵S(O)₂R¹³, -R¹⁴, -X⁵OR¹⁴, -X⁵SR¹⁴, -X⁵S(O)R¹⁴, -X⁵S(O)₂R¹⁴, -X⁵C(O)R¹⁴, -X⁵C(O)OR¹⁴, -X⁵OC(O)R¹⁴, -X⁵NR¹⁴R¹², -X⁵NR¹²C(O)R¹⁴, -X⁵NR¹²C(O)OR¹⁴, -X⁵C(O)NR¹²R¹², -X⁵S(O)₂NR¹⁴R¹², -X⁵NR¹²S(O)₂R¹⁴, -X⁵NR¹²C(O)NR¹⁴R¹² and -X⁵NR¹²C(NR¹²)NR¹⁴R¹², wherein X⁵, R¹², R¹³ and R¹⁴ are as defined above; or R¹ and R² taken together with the carbon atom to which both R¹ and R² are attached form (C₃₋₈)cycloalkylene or (C₃₋₈)heterocycloalkylene; wherein within said R² any heteroaryl, aryl, cycloalkyl, heterocycloalkyl, cycloalkylene or heterocycloalkylene is unsubstituted or substituted with 1 to 3 radicals independently selected from (C₁₋₆)alkyl, (C₁₋₆)alkylidene, cyano, halo, halo-substituted(C₁₋₄)alkyl, nitro, -X⁵NR¹²R¹², -X⁵NR¹²C(O)R¹², -X⁵NR¹²C(O)OR¹², -X⁵NR¹²C(O)NR¹²R¹², -X⁵NR¹²C(NR¹²)NR¹²R¹², -X⁵OR¹², -X⁵SR¹², -X⁵C(O)OR¹², -X⁵C(O)R¹², -X⁵OC(O)R¹², -X⁵C(O)NR¹²R¹², -X⁵S(O)₂NR¹²R¹², -X⁵NR¹²S(O)₂R¹², -X⁵P(O)(OR¹²)OR¹², -X⁵OP(O)(OR¹²)OR¹², -X⁵NR¹²C(O)R¹³, -X⁵S(O)R¹³, -X⁵S(O)₂R¹³ and -X⁵C(O)R¹³, wherein X⁵, R¹² and R¹³ are as defined above;

R³ is (C₁₋₆)alkyl or -C(R⁶)(R⁶)X⁶, wherein R⁶ is hydrogen or (C₁₋₆)alkyl and X⁶ is selected from -X⁵NR¹²R¹², -X⁵NR¹²C(O)R¹², -X⁵NR¹²C(O)OR¹², -X⁵NR¹²C(O)NR¹²R¹², -X⁵NR¹²C(NR¹²)NR¹²R¹², -X⁵OR¹², -X⁵SR¹², -X⁵C(O)OR¹², -X⁵C(O)R¹², -X⁵OC(O)R¹²,

-155-

-X⁵C(O)NR¹²R¹², -X⁵S(O)₂NR¹²R¹², -X⁵NR¹²S(O)₂R¹², -X⁵P(O)(OR¹²)OR¹²,
 -X⁵OP(O)(OR¹²)OR¹², -X⁵C(O)R¹³, -X⁵NR¹²C(O)R¹³, -X⁵S(O)R¹³, -X⁵S(O)₂R¹³, -R¹⁴,
 -X⁵OR¹⁴, -X⁵SR¹⁴, -X⁵S(O)R¹⁴, -X⁵S(O)₂R¹⁴, -X⁵C(O)R¹⁴, -X⁵C(O)OR¹⁴, -X⁵OC(O)R¹⁴,
 -X⁵NR¹⁴R¹², -X⁵NR¹²C(O)R¹⁴, -X⁵NR¹²C(O)OR¹⁴, -X⁵C(O)NR¹⁴R¹², -X⁵S(O)₂NR¹⁴R¹²,
 5 -X⁵NR¹²S(O)₂R¹⁴, -X⁵NR¹²C(O)NR¹⁴R¹² and -X⁵NR¹²C(NR¹²)NR¹⁴R¹² wherein X⁵, R¹², R¹³
 and R¹⁴ are as defined above;

R⁴ is selected from -X⁸NR¹²R¹², -X⁸NR¹²C(O)R¹², -X⁸NR¹²C(O)OR¹²,
 -X⁸NR¹²C(O)NR¹²R¹², -X⁸NR¹²C(NR¹²)NR¹²R¹², -X⁸OR¹², -X⁸SR¹², -X⁵C(O)OR¹²,
 -X⁵C(O)R¹², -X⁸OC(O)R¹², -X⁵C(O)NR¹²R¹², -X⁸S(O)₂NR¹²R¹², -X⁸NR¹²S(O)₂R¹²,
 10 -X⁸P(O)(OR¹²)OR¹², -X⁸OP(O)(OR¹²)OR¹², -X⁵C(O)R¹³, -X⁸NR¹²C(O)R¹³, -X⁸S(O)R¹³,
 -X⁸S(O)₂R¹³, -R¹⁴, -X⁸OR¹⁴, -X⁸SR¹⁴, -X⁸S(O)R¹⁴, -X⁸S(O)₂R¹⁴, -X⁵C(O)R¹⁴, -X⁵C(O)OR¹⁴,
 -X⁸OC(O)R¹⁴, -X⁸NR¹⁴R¹², -X⁸NR¹²C(O)R¹⁴, -X⁸NR¹²C(O)OR¹⁴, -X⁵C(O)NR¹⁴R¹²,
 -X⁸S(O)₂NR¹⁴R¹², -X⁸NR¹²S(O)₂R¹⁴, -X⁸NR¹²C(O)NR¹⁴R¹² and -X⁸NR¹²C(NR¹²)NR¹⁴R¹²
 wherein X⁸ is (C₁₋₆)alkylene and X⁵, R¹², R¹³ and R¹⁴ are as defined above, with the proviso
 15 that when X³ is cyano and X² is -OR⁴, where R⁴ is defined as -R¹⁴, then R¹⁴ is
 (C₃₋₁₀)cycloalkyl(C₁₋₆)alkyl, hetero(C₃₋₁₀)cycloalkyl(C₁₋₃)alkyl, (C₆₋₁₀)aryl(C₁₋₆)alkyl,
 hetero(C₅₋₁₀)aryl(C₁₋₆)alkyl, (C₉₋₁₀)bicycloaryl(C₁₋₆)alkyl or
 hetero(C₈₋₁₀)bicycloaryl(C₁₋₆)alkyl;

R¹⁵ is (C₆₋₁₀)aryl, hetero(C₅₋₁₀)aryl, (C₉₋₁₀)bicycloaryl or hetero(C₈₋₁₀)bicycloaryl;

20 R¹⁷ is (C₁₋₆)alkyl, (C₃₋₁₀)cycloalkyl(C₀₋₆)alkyl, hetero(C₃₋₁₀)cycloalkyl(C₀₋₃)alkyl,
 (C₆₋₁₀)aryl(C₀₋₆)alkyl, hetero(C₅₋₁₀)aryl(C₀₋₆)alkyl, (C₉₋₁₀)bicycloaryl(C₀₋₆)alkyl or
 hetero(C₈₋₁₀)bicycloaryl(C₀₋₆)alkyl, with the proviso that when X³ is cyano, then R¹⁷ is
 (C₁₋₆)alkyl, (C₃₋₁₀)cycloalkyl(C₁₋₆)alkyl, hetero(C₃₋₁₀)cycloalkyl(C₁₋₆)alkyl,
 (C₆₋₁₀)aryl(C₁₋₆)alkyl, hetero(C₅₋₁₀)aryl(C₁₋₆)alkyl, (C₉₋₁₀)bicycloaryl(C₁₋₆)alkyl or
 25 hetero(C₈₋₁₀)bicycloaryl(C₁₋₆)alkyl;

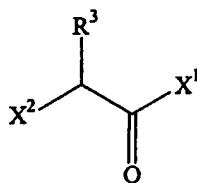
R¹⁸ is hydrogen, (C₁₋₆)alkyl, (C₃₋₁₀)cycloalkyl(C₀₋₆)alkyl,
 hetero(C₃₋₁₀)cycloalkyl(C₀₋₆)alkyl, (C₆₋₁₀)aryl(C₀₋₆)alkyl, hetero(C₅₋₁₀)aryl(C₀₋₆)alkyl,
 (C₉₋₁₀)bicycloaryl(C₀₋₆)alkyl or hetero(C₈₋₁₀)bicycloaryl(C₀₋₆)alkyl, with the proviso that when
 X³ is cyano, then R¹⁸ is (C₁₋₆)alkyl, (C₃₋₁₀)cycloalkyl(C₁₋₆)alkyl,
 30 hetero(C₃₋₁₀)cycloalkyl(C₁₋₆)alkyl, (C₆₋₁₀)aryl(C₁₋₆)alkyl, hetero(C₅₋₁₀)aryl(C₁₋₆)alkyl,
 (C₉₋₁₀)bicycloaryl(C₁₋₆)alkyl or hetero(C₈₋₁₀)bicycloaryl(C₁₋₆)alkyl; and

wherein within R³, R⁴, R¹⁵, R¹⁷ and R¹⁸ any alicyclic or aromatic ring system is
 unsubstituted or substituted further by 1 to 5 radicals independently selected from (C₁₋₆)alkyl,

-156-

(C₁₋₆)alkylidene, cyano, halo, halo-substituted(C₁₋₄)alkyl, nitro, -X⁵NR¹²R¹², -X⁵NR¹²C(O)R¹²,
 -X⁵NR¹²C(O)OR¹², -X⁵NR¹²C(O)NR¹²R¹², -X⁵NR¹²C(NR¹²)NR¹²R¹², -X⁵OR¹², -X⁵SR¹²,
 -X⁵C(O)OR¹², -X⁵C(O)R¹², -X⁵OC(O)R¹², -X⁵C(O)NR¹²R¹², -X⁵S(O)₂NR¹²R¹²,
 -X⁵NR¹²S(O)₂R¹², -X⁵P(O)(OR¹²)OR¹², -X⁵OP(O)(OR¹²)OR¹², -X⁵NR¹²C(O)R¹³, -X⁵S(O)R¹³,
 5 -X⁵C(O)R¹³ and -X⁵S(O)₂R¹³ and/or 1 radical selected from -R¹⁴, -X⁵OR¹⁴, -X⁵SR¹⁴,
 -X⁵S(O)R¹⁴, -X⁵S(O)₂R¹⁴, -X⁵C(O)R¹⁴, -X⁵C(O)OR¹⁴, -X⁵OC(O)R¹⁴, -X⁵NR¹⁴R¹²,
 -X⁵NR¹²C(O)R¹⁴, -X⁵NR¹²C(O)OR¹⁴, -X⁵C(O)NR¹⁴R¹², -X⁵S(O)₂NR¹⁴R¹², -X⁵NR¹²S(O)₂R¹⁴,
 -X⁵NR¹²C(O)NR¹⁴R¹² and -X⁵NR¹²C(NR¹²)NR¹⁴R¹²; and within R³ and R⁴ any aliphatic
 moiety is unsubstituted or substituted further by 1 to 5 radicals independently selected from
 10 cyano, halo, nitro, -NR¹²R¹², -NR¹²C(O)R¹², -NR¹²C(O)OR¹², -NR¹²C(O)NR¹²R¹²,
 -NR¹²C(NR¹²)NR¹²R¹², -OR¹², -SR¹², -C(O)OR¹², -C(O)R¹², -OC(O)R¹², -C(O)NR¹²R¹²,
 -S(O)₂NR¹²R¹², -NR¹²S(O)₂R¹², -P(O)(OR¹²)OR¹², -OP(O)(OR¹²)OR¹², -NR¹²C(O)R¹³,
 -S(O)R¹³ and -S(O)₂R¹³; wherein X⁵, R¹², R¹³ and R¹⁴ are as described above, with the proviso
 that when X³ is cyano and X² is -OR⁴, where R⁴ is defined as -R¹⁴, or -NHR¹⁸, then any
 15 aromatic ring system present within R¹⁴ or R¹⁸ is not substituted further by halo,
 (C₃₋₁₀)cycloalkyl, hetero(C₃₋₁₀)cycloalkyl, (C₆₋₁₀)aryl, hetero(C₅₋₁₀)aryl, (C₉₋₁₀)bicycloaryl or
 hetero(C₈₋₁₀)bicycloaryl; with the proviso that only one bicyclic ring structure is present
 within R³, R⁴ or R¹⁵; and the *N*-oxide derivatives, prodrug derivatives, protected derivatives,
 individual isomers and mixtures of isomers thereof; and the pharmaceutically acceptable salts
 20 and solvates of such compounds and the *N*-oxide derivatives, prodrug derivatives, protected
 derivatives, individual isomers and mixtures of isomers thereof.

2. A compound of Claim 1, which is of the following formula:



25

in which X² is hydrogen, fluoro, -OH, -OR⁴, -NHR¹⁵;
 R³, R⁴, R¹⁵ and X¹ are the same as defined in claim 1.

30

3. A compound of Claim 1 or Claim 2 in which:

-157-

X^1 is $-\text{NHC}(\text{R}^1)(\text{R}^2)\text{X}^3$ or $-\text{NHCH}(\text{R}^{19})\text{C}(\text{O})\text{R}^{20}$;

X^2 is hydrogen, fluoro, $-\text{OH}$, $-\text{OR}^4$, $-\text{NHR}^{15}$ or $-\text{NR}^{17}\text{R}^{18}$ and X^7 is hydrogen or X^2 and X^7 both represent fluoro;

X^3 is cyano, $-\text{C}(\text{R}^7)(\text{R}^8)\text{R}^{16}$, $-\text{C}(\text{R}^6)(\text{OR}^6)_2$, $-\text{CH}_2\text{C}(\text{O})\text{R}^{16}$, $-\text{CH}=\text{CHS}(\text{O})_2\text{R}^5$,
 5 $-\text{C}(\text{O})\text{CF}_2\text{C}(\text{O})\text{NR}^5\text{R}^5$, $-\text{C}(\text{O})\text{C}(\text{O})\text{NR}^5\text{R}^6$, $-\text{C}(\text{O})\text{C}(\text{O})\text{OR}^5$, $-\text{C}(\text{O})\text{CH}_2\text{OR}^5$,
 $-\text{C}(\text{O})\text{CH}_2\text{N}(\text{R}^6)\text{SO}_2\text{R}^5$ or $-\text{C}(\text{O})\text{C}(\text{O})\text{R}^5$; wherein R^5 is hydrogen, (C_{1-4}) alkyl,
 (C_{3-10}) cycloalkyl (C_{0-6}) alkyl, hetero (C_{3-10}) cycloalkyl (C_{0-3}) alkyl, (C_{6-10}) aryl (C_{0-6}) alkyl,
 hetero (C_{5-10}) aryl (C_{0-6}) alkyl, (C_{9-10}) bicycloaryl (C_{0-6}) alkyl or
 hetero (C_{8-10}) bicycloaryl (C_{0-6}) alkyl; R^6 is hydrogen, hydroxy or (C_{1-6}) alkyl; or where X^3
 10 contains an $-\text{NR}^5\text{R}^6$ group, R^5 and R^6 together with the nitrogen atom to which they are both
 attached, form hetero (C_{3-10}) cycloalkyl, hetero (C_{5-10}) aryl or hetero (C_{8-10}) bicycloaryl; R^7 is
 hydrogen or (C_{1-4}) alkyl and R^8 is hydroxy or R^7 and R^8 together form oxo; R^{16} is hydrogen, $-\text{X}^4$,
 $-\text{CF}_3$, $-\text{CF}_2\text{CF}_2\text{R}^9$ or $-\text{N}(\text{R}^6)\text{OR}^6$; R^9 is hydrogen, halo, (C_{1-4}) alkyl, (C_{5-10}) aryl (C_{0-6}) alkyl or
 (C_{5-10}) heteroaryl (C_{0-6}) alkyl, with the proviso that when X^3 is cyano, then X^2 is hydrogen,
 15 fluoro, $-\text{OH}$, $-\text{OR}^4$ or $-\text{NR}^{17}\text{R}^{18}$ and X^7 is hydrogen or X^2 and X^7 both represent fluoro;

X^4 comprises a heteromonocyclic ring containing 4 to 7 ring member atoms or a fused
 heterobicyclic ring system containing 8 to 14 ring member atoms and any carbocyclic ketone,
 iminoketone or thioketone derivative thereof, with the proviso that when $-\text{X}^4$ is other than a
 heteromonocyclic ring containing 5 ring member atoms, wherein no more than two of the ring
 20 member atoms comprising the ring are heteroatoms, then X^2 is fluoro, $-\text{OH}$, $-\text{OR}^4$, $-\text{NHR}^{15}$ or
 $-\text{NR}^{17}\text{R}^{18}$ and X^7 is hydrogen or X^2 and X^7 both represent fluoro;

wherein within R^5 , X^3 or X^4 any alicyclic or aromatic ring system is unsubstituted or
 substituted further by 1 to 5 radicals independently selected from (C_{1-6}) alkyl, (C_{1-6}) alkylidene,
 cyano, halo, halo-substituted (C_{1-4}) alkyl, nitro, $-\text{X}^5\text{NR}^{12}\text{R}^{12}$, $-\text{X}^5\text{NR}^{12}\text{C}(\text{O})\text{R}^{12}$,
 25 $-\text{X}^5\text{NR}^{12}\text{C}(\text{O})\text{OR}^{12}$, $-\text{X}^5\text{NR}^{12}\text{C}(\text{O})\text{NR}^{12}\text{R}^{12}$, $-\text{X}^5\text{NR}^{12}\text{C}(\text{NR}^{12})\text{NR}^{12}\text{R}^{12}$, $-\text{X}^5\text{OR}^{12}$, $-\text{X}^5\text{SR}^{12}$,
 $-\text{X}^5\text{C}(\text{O})\text{OR}^{12}$, $-\text{X}^5\text{C}(\text{O})\text{R}^{12}$, $-\text{X}^5\text{OC}(\text{O})\text{R}^{12}$, $-\text{X}^5\text{C}(\text{O})\text{NR}^{12}\text{R}^{12}$, $-\text{X}^5\text{S}(\text{O})_2\text{NR}^{12}\text{R}^{12}$,
 $-\text{X}^5\text{NR}^{12}\text{S}(\text{O})_2\text{R}^{12}$, $-\text{X}^5\text{P}(\text{O})(\text{OR}^{12})\text{OR}^{12}$, $-\text{X}^5\text{OP}(\text{O})(\text{OR}^{12})\text{OR}^{12}$, $-\text{X}^5\text{NR}^{12}\text{C}(\text{O})\text{R}^{13}$, $-\text{X}^5\text{S}(\text{O})\text{R}^{13}$
 and $-\text{X}^5\text{S}(\text{O})_2\text{R}^{13}$ and/or 1 radical selected from $-\text{R}^{14}$, $-\text{X}^5\text{OR}^{14}$, $-\text{X}^5\text{SR}^{14}$, $-\text{X}^5\text{S}(\text{O})\text{R}^{14}$,
 $-\text{X}^5\text{S}(\text{O})_2\text{R}^{14}$, $-\text{X}^5\text{C}(\text{O})\text{R}^{14}$, $-\text{X}^5\text{C}(\text{O})\text{OR}^{14}$, $-\text{X}^5\text{OC}(\text{O})\text{R}^{14}$, $-\text{X}^5\text{NR}^{14}\text{R}^{12}$, $-\text{X}^5\text{NR}^{12}\text{C}(\text{O})\text{R}^{14}$,
 30 $-\text{X}^5\text{NR}^{12}\text{C}(\text{O})\text{OR}^{14}$, $-\text{X}^5\text{C}(\text{O})\text{NR}^{12}\text{R}^{12}$, $-\text{X}^5\text{S}(\text{O})_2\text{NR}^{14}\text{R}^{12}$, $-\text{X}^5\text{NR}^{12}\text{S}(\text{O})_2\text{R}^{14}$,
 $-\text{X}^5\text{NR}^{12}\text{C}(\text{O})\text{NR}^{14}\text{R}^{12}$ and $-\text{X}^5\text{NR}^{12}\text{C}(\text{NR}^{12})\text{NR}^{14}\text{R}^{12}$, wherein X^5 is a bond or (C_{1-6}) alkylene;
 R^{12} at each occurrence independently is hydrogen, (C_{1-6}) alkyl or halo-substituted (C_{1-6}) alkyl;
 R^{13} is (C_{1-6}) alkyl or halo-substituted (C_{1-6}) alkyl; and R^{14} is (C_{3-10}) cycloalkyl (C_{0-6}) alkyl,

hetero(C₃₋₁₀)cycloalkyl(C₀₋₃)alkyl, (C₆₋₁₀)aryl(C₀₋₆)alkyl, hetero(C₅₋₁₀)aryl(C₀₋₆)alkyl, (C₉₋₁₀)bicycloaryl(C₀₋₆)alkyl or hetero(C₈₋₁₀)bicycloaryl(C₀₋₆)alkyl;

R¹ is hydrogen or (C₁₋₆)alkyl and R² is selected from a group consisting of hydrogen, cyano, -X⁵NR¹²R¹², -X⁵NR¹²C(O)R¹², -X⁵NR¹²C(O)OR¹², -R¹², -X⁵NR¹²C(O)NR¹²R¹²,
 5 -X⁵NR¹²C(NR¹²)NR¹²R¹², -X⁵OR¹², -X⁵SR¹², -X⁵C(O)OR¹², -X⁵C(O)R¹², -X⁵OC(O)R¹²,
 -X⁵C(O)NR¹²R¹², -X⁵S(O)₂NR¹²R¹², -X⁵NR¹²S(O)₂R¹², -X⁵P(O)(OR¹²)OR¹²,
 -X⁵OP(O)(OR¹²)OR¹², -X⁵NR¹²C(O)R¹³, -X⁵S(O)R¹³, -X⁵S(O)₂R¹³, -R¹⁴, -X⁵OR¹⁴, -X⁵SR¹⁴,
 -X⁵S(O)R¹⁴, -X⁵S(O)₂R¹⁴, -X⁵C(O)R¹⁴, -X⁵C(O)OR¹⁴, -X⁵OC(O)R¹⁴, -X⁵NR¹⁴R¹²,
 -X⁵NR¹²C(O)R¹⁴, -X⁵NR¹²C(O)OR¹⁴, -X⁵C(O)NR¹²R¹², -X⁵S(O)₂NR¹⁴R¹², -X⁵NR¹²S(O)₂R¹⁴,
 10 -X⁵NR¹²C(O)NR¹⁴R¹² and -X⁵NR¹²C(NR¹²)NR¹⁴R¹², wherein X⁵, R¹², R¹³ and R¹⁴ are as defined above; or R¹ and R² taken together with the carbon atom to which both R¹ and R² are attached form (C₃₋₈)cycloalkylene or (C₃₋₈)heterocycloalkylene; wherein within said R² any heteroaryl, aryl, cycloalkyl, heterocycloalkyl, cycloalkylene or heterocycloalkylene is unsubstituted or substituted with 1 to 3 radicals independently selected from (C₁₋₆)alkyl,
 15 (C₁₋₆)alkylidene, cyano, halo, halo-substituted(C₁₋₄)alkyl, nitro, -X⁵NR¹²R¹², -X⁵NR¹²C(O)R¹²,
 -X⁵NR¹²C(O)OR¹², -X⁵NR¹²C(O)NR¹²R¹², -X⁵NR¹²C(NR¹²)NR¹²R¹², -X⁵OR¹², -X⁵SR¹²,
 -X⁵C(O)OR¹², -X⁵C(O)R¹², -X⁵OC(O)R¹², -X⁵C(O)NR¹²R¹², -X⁵S(O)₂NR¹²R¹²,
 -X⁵NR¹²S(O)₂R¹², -X⁵P(O)(OR¹²)OR¹², -X⁵OP(O)(OR¹²)OR¹², -X⁵NR¹²C(O)R¹³, -X⁵S(O)R¹³,
 -X⁵S(O)₂R¹³ and -X⁵C(O)R¹³, wherein X⁵, R¹² and R¹³ are as defined above;

20 R³ is (C₁₋₆)alkyl or -C(R⁶)(R⁶)X⁶, wherein R⁶ is hydrogen or (C₁₋₆)alkyl and X⁶ is selected from -X⁵NR¹²R¹², -X⁵NR¹²C(O)R¹², -X⁵NR¹²C(O)OR¹², -X⁵NR¹²C(O)NR¹²R¹²,
 -X⁵NR¹²C(NR¹²)NR¹²R¹², -X⁵OR¹², -X⁵SR¹², -X⁵C(O)OR¹², -X⁵C(O)R¹², -X⁵OC(O)R¹²,
 -X⁵C(O)NR¹²R¹², -X⁵S(O)₂NR¹²R¹², -X⁵NR¹²S(O)₂R¹², -X⁵P(O)(OR¹²)OR¹²,
 -X⁵OP(O)(OR¹²)OR¹², -X⁵C(O)R¹³, -X⁵NR¹²C(O)R¹³, -X⁵S(O)R¹³, -X⁵S(O)₂R¹³, -R¹⁴,
 25 -X⁵OR¹⁴, -X⁵SR¹⁴, -X⁵S(O)R¹⁴, -X⁵S(O)₂R¹⁴, -X⁵C(O)R¹⁴, -X⁵C(O)OR¹⁴, -X⁵OC(O)R¹⁴,
 -X⁵NR¹⁴R¹², -X⁵NR¹²C(O)R¹⁴, -X⁵NR¹²C(O)OR¹⁴, -X⁵C(O)NR¹⁴R¹², -X⁵S(O)₂NR¹⁴R¹²,
 -X⁵NR¹²S(O)₂R¹⁴, -X⁵NR¹²C(O)NR¹⁴R¹² and -X⁵NR¹²C(NR¹²)NR¹⁴R¹² wherein X⁵, R¹², R¹³ and R¹⁴ are as defined above;

R⁴ is selected from -X⁸NR¹²R¹², -X⁸NR¹²C(O)R¹², -X⁸NR¹²C(O)OR¹²,
 30 -X⁸NR¹²C(O)NR¹²R¹², -X⁸NR¹²C(NR¹²)NR¹²R¹², -X⁸OR¹², -X⁸SR¹², -X⁵C(O)OR¹²,
 -X⁵C(O)R¹², -X⁸OC(O)R¹², -X⁵C(O)NR¹²R¹², -X⁸S(O)₂NR¹²R¹², -X⁸NR¹²S(O)₂R¹²,
 -X⁸P(O)(OR¹²)OR¹², -X⁸OP(O)(OR¹²)OR¹², -X⁵C(O)R¹³, -X⁸NR¹²C(O)R¹³, -X⁸S(O)R¹³,
 -X⁸S(O)₂R¹³, -R¹⁴, -X⁸OR¹⁴, -X⁸SR¹⁴, -X⁸S(O)R¹⁴, -X⁸S(O)₂R¹⁴, -X⁵C(O)R¹⁴, -X⁵C(O)OR¹⁴,

-159-

$-X^8OC(O)R^{14}$, $-X^8NR^{14}R^{12}$, $-X^8NR^{12}C(O)R^{14}$, $-X^8NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{14}R^{12}$,
 $-X^8S(O)_2NR^{14}R^{12}$, $-X^8NR^{12}S(O)_2R^{14}$, $-X^8NR^{12}C(O)NR^{14}R^{12}$ and $-X^8NR^{12}C(NR^{12})NR^{14}R^{12}$
 wherein X^8 is (C_{1-6}) alkylene and X^5 , R^{12} , R^{13} and R^{14} are as defined above, with the proviso
 that when X^3 is cyano and X^2 is $-OR^4$, where R^4 is defined as $-R^{14}$, then R^{14} is

- 5 (C_{3-10}) cycloalkyl (C_{1-6}) alkyl, hetero (C_{3-10}) cycloalkyl (C_{1-3}) alkyl, (C_{6-10}) aryl (C_{1-6}) alkyl,
 hetero (C_{5-10}) aryl (C_{1-6}) alkyl, (C_{9-10}) bicycloaryl (C_{1-6}) alkyl or
 hetero (C_{8-10}) bicycloaryl (C_{1-6}) alkyl;

R^{15} is (C_{6-10}) aryl, hetero (C_{5-10}) aryl, (C_{9-10}) bicycloaryl or hetero (C_{8-10}) bicycloaryl;

- R^{17} is (C_{1-6}) alkyl, (C_{3-10}) cycloalkyl (C_{0-6}) alkyl, hetero (C_{3-10}) cycloalkyl (C_{0-3}) alkyl,
 10 (C_{6-10}) aryl (C_{0-6}) alkyl, hetero (C_{5-10}) aryl (C_{0-6}) alkyl, (C_{9-10}) bicycloaryl (C_{0-6}) alkyl or
 hetero (C_{8-10}) bicycloaryl (C_{0-6}) alkyl, with the proviso that when X^3 is cyano, then R^{17} is
 (C_{1-6}) alkyl, (C_{3-10}) cycloalkyl (C_{1-6}) alkyl, hetero (C_{3-10}) cycloalkyl (C_{1-6}) alkyl,
 (C_{6-10}) aryl (C_{1-6}) alkyl, hetero (C_{5-10}) aryl (C_{1-6}) alkyl, (C_{9-10}) bicycloaryl (C_{1-6}) alkyl or
 hetero (C_{8-10}) bicycloaryl (C_{1-6}) alkyl;

- 15 R^{18} is hydrogen, (C_{1-6}) alkyl, (C_{3-10}) cycloalkyl (C_{0-6}) alkyl,
 hetero (C_{3-10}) cycloalkyl (C_{0-6}) alkyl, (C_{6-10}) aryl (C_{0-6}) alkyl, hetero (C_{5-10}) aryl (C_{0-6}) alkyl,
 (C_{9-10}) bicycloaryl (C_{0-6}) alkyl or hetero (C_{8-10}) bicycloaryl (C_{0-6}) alkyl, with the proviso that when
 X^3 is cyano, then R^{18} is (C_{1-6}) alkyl, (C_{3-10}) cycloalkyl (C_{1-6}) alkyl,
 hetero (C_{3-10}) cycloalkyl (C_{1-6}) alkyl, (C_{6-10}) aryl (C_{1-6}) alkyl, hetero (C_{5-10}) aryl (C_{1-6}) alkyl,
 20 (C_{9-10}) bicycloaryl (C_{1-6}) alkyl or hetero (C_{8-10}) bicycloaryl (C_{1-6}) alkyl; and

- R^{19} and R^{20} together with the atoms to which R^{19} and R^{20} are attached form
 (C_{4-8}) heterocycloalkylene, wherein no more than one of the ring member atoms comprising
 the ring is a heteroatom selected from $-NR^{21}-$ or $-O-$, wherein the ring is unsubstituted or
 substituted with R^2 , wherein R^2 is as defined above, and R^{21} is hydrogen, $-C(O)OR^{12}$,
 25 $-C(O)R^{12}$, $-C(O)NR^{12}R^{12}$, $-S(O)_2NR^{12}R^{12}$, $-S(O)R^{13}$ and $-S(O)_2R^{13}$, $-S(O)R^{14}$, $-S(O)_2R^{14}$,
 $-C(O)R^{14}$, $-C(O)OR^{14}$, $-C(O)NR^{12}R^{12}$ and $-S(O)_2NR^{14}R^{12}$, wherein R^{12} , R^{13} and R^{14} are as
 defined above;

- wherein within R^3 , R^4 , R^{15} , R^{17} and R^{18} any alicyclic or aromatic ring system is
 unsubstituted or substituted further by 1 to 5 radicals independently selected from (C_{1-6}) alkyl,
 30 (C_{1-6}) alkylidene, cyano, halo, halo-substituted (C_{1-4}) alkyl, nitro, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$,
 $-X^5NR^{12}C(O)OR^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$,
 $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$,
 $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$,

-160-

$-X^5C(O)R^{13}$ and $-X^5S(O)_2R^{13}$ and/or 1 radical selected from $-R^{14}$, $-X^5OR^{14}$, $-X^5SR^{14}$,
 $-X^5S(O)R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$, $-X^5OC(O)R^{14}$, $-X^5NR^{14}R^{12}$,
 $-X^5NR^{12}C(O)R^{14}$, $-X^5NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{14}R^{12}$, $-X^5S(O)_2NR^{14}R^{12}$, $-X^5NR^{12}S(O)_2R^{14}$,
 $-X^5NR^{12}C(O)NR^{14}R^{12}$ and $-X^5NR^{12}C(NR^{12})NR^{14}R^{12}$; and within R^3 and R^4 any aliphatic
5 moiety is unsubstituted or substituted further by 1 to 5 radicals independently selected from
cyano, halo, nitro, $-NR^{12}R^{12}$, $-NR^{12}C(O)R^{12}$, $-NR^{12}C(O)OR^{12}$, $-NR^{12}C(O)NR^{12}R^{12}$,
 $-NR^{12}C(NR^{12})NR^{12}R^{12}$, $-OR^{12}$, $-SR^{12}$, $-C(O)OR^{12}$, $-C(O)R^{12}$, $-OC(O)R^{12}$, $-C(O)NR^{12}R^{12}$,
 $-S(O)_2NR^{12}R^{12}$, $-NR^{12}S(O)_2R^{12}$, $-P(O)(OR^{12})OR^{12}$, $-OP(O)(OR^{12})OR^{12}$, $-NR^{12}C(O)R^{13}$,
 $-S(O)R^{13}$ and $-S(O)_2R^{13}$; wherein X^5 , R^{12} , R^{13} and R^{14} are as described above, with the proviso
10 that when X^3 is cyano and X^2 is $-OR^4$, where R^4 is defined as $-R^{14}$, or $-NHR^{18}$, then any
aromatic ring system present within R^{14} or R^{18} is not substituted further by halo,
 (C_{3-10}) cycloalkyl, hetero (C_{3-10}) cycloalkyl, (C_{6-10}) aryl, hetero (C_{5-10}) aryl, (C_{9-10}) bicycloaryl or
hetero (C_{8-10}) bicycloaryl; with the proviso that only one bicyclic ring structure is present
within R^3 , R^4 or R^{15} ; and the *N*-oxide derivatives, prodrug derivatives, protected derivatives,
15 individual isomers and mixtures of isomers thereof; and the pharmaceutically acceptable salts
and solvates of such compounds and the *N*-oxide derivatives, prodrug derivatives, protected
derivatives, individual isomers and mixtures of isomers thereof.

4. The compound of Claim 1 or Claim 2 in which:
20 X^1 is $-NHC(R^1)(R^2)X^3$ or $-NHCH(R^{19})C(O)R^{20}$;
 X^2 is hydrogen, fluoro, $-OH$, $-OR^4$, $-NHR^{15}$ or $-NR^{17}R^{18}$ and X^7 is hydrogen or X^2 and
 X^7 both represent fluoro;
 X^3 is $-C(R^7)(R^8)R^{16}$, $-C(R^6)(OR^6)_2$, $-CH_2C(O)R^{16}$, $-CH=CHS(O)_2R^5$,
 $-C(O)CF_2C(O)NR^5R^5$, $-C(O)C(O)NR^5R^6$, $-C(O)C(O)OR^5$, $-C(O)CH_2OR^5$,
25 $-C(O)CH_2N(R^6)SO_2R^5$ or $-C(O)C(O)R^5$; wherein R^5 is hydrogen, (C_{1-4}) alkyl,
 (C_{3-10}) cycloalkyl (C_{0-6}) alkyl, hetero (C_{3-10}) cycloalkyl (C_{0-3}) alkyl, (C_{6-10}) aryl (C_{0-6}) alkyl,
hetero (C_{5-10}) aryl (C_{0-6}) alkyl, (C_{9-10}) bicycloaryl (C_{0-6}) alkyl or
hetero (C_{8-10}) bicycloaryl (C_{0-6}) alkyl; R^6 is hydrogen, hydroxy or (C_{1-6}) alkyl; or where X^3
contains an $-NR^5R^6$ group, R^5 and R^6 together with the nitrogen atom to which they are both
30 attached, form hetero (C_{3-10}) cycloalkyl, hetero (C_{5-10}) aryl or hetero (C_{8-10}) bicycloaryl; R^7 is
hydrogen or (C_{1-4}) alkyl and R^8 is hydroxy or R^7 and R^8 together form oxo; R^{16} is hydrogen, -
 X^4 , $-CF_3$, $-CF_2CF_2R^9$ or $-N(R^6)OR^6$; R^9 is hydrogen, halo, (C_{1-4}) alkyl, (C_{5-10}) aryl (C_{0-6}) alkyl or
 (C_{5-10}) heteroaryl (C_{0-6}) alkyl;

-161-

X^4 comprises a heteromonocyclic ring containing 4 to 7 ring member atoms or a fused heterobicyclic ring system containing 8 to 14 ring member atoms and any carbocyclic ketone, iminoketone or thioketone derivative thereof, with the proviso that when $-X^4$ is other than a heteromonocyclic ring containing 5 ring member atoms, wherein no more than two of the ring member atoms comprising the ring are heteroatoms, then X^2 is fluoro, -OH, $-OR^4$, $-NHR^{15}$ or $-NR^{17}R^{18}$ and X^7 is hydrogen or X^2 and X^7 both represent fluoro;

wherein within R^5 , X^3 or X^4 any alicyclic or aromatic ring system is unsubstituted or substituted further by 1 to 5 radicals independently selected from (C_{1-6}) alkyl, (C_{1-6}) alkylidene, cyano, halo, halo-substituted (C_{1-4}) alkyl, nitro, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$, $-X^5NR^{12}C(O)OR^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$, $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$, $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$ and $-X^5S(O)_2R^{13}$ and/or 1 radical selected from $-R^{14}$, $-X^5OR^{14}$, $-X^5SR^{14}$, $-X^5S(O)R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$, $-X^5OC(O)R^{14}$, $-X^5NR^{14}R^{12}$, $-X^5NR^{12}C(O)R^{14}$, $-X^5NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{14}R^{12}$, $-X^5NR^{12}S(O)_2R^{14}$, $-X^5NR^{12}C(O)NR^{14}R^{12}$ and $-X^5NR^{12}C(NR^{12})NR^{14}R^{12}$, wherein X^5 is a bond or (C_{1-6}) alkylene; R^{12} at each occurrence independently is hydrogen, (C_{1-6}) alkyl or halo-substituted (C_{1-6}) alkyl; R^{13} is (C_{1-6}) alkyl or halo-substituted (C_{1-6}) alkyl; and R^{14} is (C_{3-10}) cycloalkyl (C_{0-6}) alkyl, hetero (C_{3-10}) cycloalkyl (C_{0-3}) alkyl, (C_{6-10}) aryl (C_{0-6}) alkyl, hetero (C_{5-10}) aryl (C_{0-6}) alkyl, (C_{9-10}) bicycloaryl (C_{0-6}) alkyl or hetero (C_{8-10}) bicycloaryl (C_{0-6}) alkyl;

R^1 is hydrogen or (C_{1-6}) alkyl and R^2 is selected from a group consisting of hydrogen, cyano, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$, $-X^5NR^{12}C(O)OR^{12}$, $-R^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$, $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$, $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$, $-X^5S(O)_2R^{13}$, $-R^{14}$, $-X^5OR^{14}$, $-X^5SR^{14}$, $-X^5S(O)R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$, $-X^5OC(O)R^{14}$, $-X^5NR^{14}R^{12}$, $-X^5NR^{12}C(O)R^{14}$, $-X^5NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{14}R^{12}$, $-X^5NR^{12}S(O)_2R^{14}$, $-X^5NR^{12}C(O)NR^{14}R^{12}$ and $-X^5NR^{12}C(NR^{12})NR^{14}R^{12}$, wherein X^5 , R^{12} , R^{13} and R^{14} are as defined above; or R^1 and R^2 taken together with the carbon atom to which both R^1 and R^2 are attached form (C_{3-8}) cycloalkylene or (C_{3-8}) heterocycloalkylene; wherein within said R^2 any heteroaryl, aryl, cycloalkyl, heterocycloalkyl, cycloalkylene or heterocycloalkylene is unsubstituted or substituted with 1 to 3 radicals independently selected from (C_{1-6}) alkyl, (C_{1-6}) alkylidene, cyano, halo, halo-substituted (C_{1-4}) alkyl, nitro, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$,

-162-

$-X^5NR^{12}C(O)OR^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$,
 $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$,
 $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$,
 $-X^5S(O)_2R^{13}$ and $-X^5C(O)R^{13}$, wherein X^5 , R^{12} and R^{13} are as defined above;

- 5 R^3 is (C_{1-6}) alkyl or $-C(R^6)(R^6)X^6$, wherein R^6 is hydrogen or (C_{1-6}) alkyl and X^6 is selected from $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$, $-X^5NR^{12}C(O)OR^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$, $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$, $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5C(O)R^{13}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$, $-X^5S(O)_2R^{13}$, $-R^{14}$,
 10 $-X^5OR^{14}$, $-X^5SR^{14}$, $-X^5S(O)R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$, $-X^5OC(O)R^{14}$, $-X^5NR^{14}R^{12}$, $-X^5NR^{12}C(O)R^{14}$, $-X^5NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{14}R^{12}$, $-X^5S(O)_2NR^{14}R^{12}$, $-X^5NR^{12}S(O)_2R^{14}$, $-X^5NR^{12}C(O)NR^{14}R^{12}$ and $-X^5NR^{12}C(NR^{12})NR^{14}R^{12}$ wherein X^5 , R^{12} , R^{13} and R^{14} are as defined above;

- R^4 is selected from $-X^8NR^{12}R^{12}$, $-X^8NR^{12}C(O)R^{12}$, $-X^8NR^{12}C(O)OR^{12}$,
 15 $-X^8NR^{12}C(O)NR^{12}R^{12}$, $-X^8NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^8OR^{12}$, $-X^8SR^{12}$, $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^8OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^8S(O)_2NR^{12}R^{12}$, $-X^8NR^{12}S(O)_2R^{12}$, $-X^8P(O)(OR^{12})OR^{12}$, $-X^8OP(O)(OR^{12})OR^{12}$, $-X^5C(O)R^{13}$, $-X^8NR^{12}C(O)R^{13}$, $-X^8S(O)R^{13}$, $-X^8S(O)_2R^{13}$, $-R^{14}$, $-X^8OR^{14}$, $-X^8SR^{14}$, $-X^8S(O)R^{14}$, $-X^8S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$, $-X^8OC(O)R^{14}$, $-X^8NR^{14}R^{12}$, $-X^8NR^{12}C(O)R^{14}$, $-X^8NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{14}R^{12}$,
 20 $-X^8S(O)_2NR^{14}R^{12}$, $-X^8NR^{12}S(O)_2R^{14}$, $-X^8NR^{12}C(O)NR^{14}R^{12}$ and $-X^8NR^{12}C(NR^{12})NR^{14}R^{12}$ wherein X^8 is (C_{1-6}) alkylene and X^5 , R^{12} , R^{13} and R^{14} are as defined above;

R^{15} is (C_{6-10}) aryl, hetero (C_{5-10}) aryl, (C_{9-10}) bicycloaryl or hetero (C_{8-10}) bicycloaryl;

- R^{17} is hydrogen, (C_{1-6}) alkyl, (C_{3-10}) cycloalkyl (C_{0-6}) alkyl,
 hetero (C_{3-10}) cycloalkyl (C_{0-6}) alkyl, (C_{6-10}) aryl (C_{0-6}) alkyl, hetero (C_{5-10}) aryl (C_{0-6}) alkyl,
 25 (C_{9-10}) bicycloaryl (C_{0-6}) alkyl or hetero (C_{8-10}) bicycloaryl (C_{0-6}) alkyl;

R^{18} is (C_{1-6}) alkyl, (C_{3-10}) cycloalkyl (C_{0-6}) alkyl, hetero (C_{3-10}) cycloalkyl (C_{0-6}) alkyl,
 (C_{6-10}) aryl (C_{0-6}) alkyl, hetero (C_{5-10}) aryl (C_{0-6}) alkyl, (C_{9-10}) bicycloaryl (C_{0-6}) alkyl or
 hetero (C_{8-10}) bicycloaryl (C_{0-6}) alkyl; and

- R^{19} and R^{20} together with the atoms to which R^{19} and R^{20} are attached form
 30 (C_{4-8}) heterocycloalkylene, wherein no more than one of the ring member atoms comprising the ring is a heteroatom selected from $-NR^{21}-$ or $-O-$, wherein the ring is unsubstituted or substituted with R^2 , wherein R^2 is as defined above, and R^{21} is hydrogen, $-C(O)OR^{12}$, $-C(O)R^{12}$, $-C(O)NR^{12}R^{12}$, $-S(O)_2NR^{12}R^{12}$, $-S(O)R^{13}$ and $-S(O)_2R^{13}$, $-S(O)R^{14}$, $-S(O)_2R^{14}$,

-163-

$-C(O)R^{14}$, $-C(O)OR^{14}$, $-C(O)NR^{12}R^{12}$ and $-S(O)_2NR^{14}R^{12}$, wherein R^{12} , R^{13} and R^{14} are as defined above;

wherein within R^3 , R^4 , R^{15} , R^{17} and R^{18} any alicyclic or aromatic ring system is unsubstituted or substituted further by 1 to 5 radicals independently selected from (C_{1-6}) alkyl, (C_{1-6}) alkylidene, cyano, halo, halo-substituted (C_{1-4}) alkyl, nitro, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$, $-X^5NR^{12}C(O)OR^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$, $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$, $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$, $-X^5C(O)R^{13}$ and $-X^5S(O)_2R^{13}$ and/or 1 radical selected from $-R^{14}$, $-X^5OR^{14}$, $-X^5SR^{14}$, $-X^5S(O)R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$, $-X^5OC(O)R^{14}$, $-X^5NR^{14}R^{12}$, $-X^5NR^{12}C(O)R^{14}$, $-X^5NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{14}R^{12}$, $-X^5S(O)_2NR^{14}R^{12}$, $-X^5NR^{12}S(O)_2R^{14}$, $-X^5NR^{12}C(O)NR^{14}R^{12}$ and $-X^5NR^{12}C(NR^{12})NR^{14}R^{12}$; and within R^3 and R^4 any aliphatic moiety is unsubstituted or substituted further by 1 to 5 radicals independently selected from cyano, halo, nitro, $-NR^{12}R^{12}$, $-NR^{12}C(O)R^{12}$, $-NR^{12}C(O)OR^{12}$, $-NR^{12}C(O)NR^{12}R^{12}$, $-NR^{12}C(NR^{12})NR^{12}R^{12}$, $-OR^{12}$, $-SR^{12}$, $-C(O)OR^{12}$, $-C(O)R^{12}$, $-OC(O)R^{12}$, $-C(O)NR^{12}R^{12}$, $-S(O)_2NR^{12}R^{12}$, $-NR^{12}S(O)_2R^{12}$, $-P(O)(OR^{12})OR^{12}$, $-OP(O)(OR^{12})OR^{12}$, $-NR^{12}C(O)R^{13}$, $-S(O)R^{13}$ and $-S(O)_2R^{13}$; wherein X^5 , R^{12} , R^{13} and R^{14} are as described above; with the proviso that only one bicyclic ring structure is present within R^3 , R^4 or R^{15} ; and the *N*-oxide derivatives, prodrug derivatives, protected derivatives, individual isomers and mixtures of isomers thereof; and the pharmaceutically acceptable salts and solvates of such compounds and the *N*-oxide derivatives, prodrug derivatives, protected derivatives, individual isomers and mixtures of isomers thereof.

5. A compound of Claim 1 or Claim 2 in which:
 X^1 is $-NHC(R^1)(R^2)X^3$ or $-NHCH(R^{19})C(O)R^{20}$;
 X^2 is hydrogen, fluoro, $-OH$, $-OR^4$ or $-NR^{17}R^{18}$ and X^7 is hydrogen or X^2 and X^7 both represent fluoro;

X^3 is cyano;

wherein within X^3 any alicyclic or aromatic ring system is unsubstituted or substituted further by 1 to 5 radicals independently selected from (C_{1-6}) alkyl, (C_{1-6}) alkylidene, cyano, halo, halo-substituted (C_{1-4}) alkyl, nitro, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$, $-X^5NR^{12}C(O)OR^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$, $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$, $-X^5NR^{12}S(O)_2R^{12}$,

-164-

- $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$ and $-X^5S(O)_2R^{13}$
 and/or 1 radical selected from $-R^{14}$, $-X^5OR^{14}$, $-X^5SR^{14}$, $-X^5S(O)R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$,
 $-X^5C(O)OR^{14}$, $-X^5OC(O)R^{14}$, $-X^5NR^{14}R^{12}$, $-X^5NR^{12}C(O)R^{14}$, $-X^5NR^{12}C(O)OR^{14}$,
 $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{14}R^{12}$, $-X^5NR^{12}S(O)_2R^{14}$, $-X^5NR^{12}C(O)NR^{14}R^{12}$ and
 5 $-X^5NR^{12}C(NR^{12})NR^{14}R^{12}$, wherein X^5 is a bond or (C_{1-6}) alkylene; R^{12} at each occurrence
 independently is hydrogen, (C_{1-6}) alkyl or halo-substituted (C_{1-6}) alkyl; R^{13} is (C_{1-6}) alkyl or
 halo-substituted (C_{1-6}) alkyl; and R^{14} is (C_{3-10}) cycloalkyl (C_{0-6}) alkyl,
 hetero (C_{3-10}) cycloalkyl (C_{0-3}) alkyl, (C_{6-10}) aryl (C_{0-6}) alkyl, hetero (C_{5-10}) aryl (C_{0-6}) alkyl,
 (C_{9-10}) bicycloaryl (C_{0-6}) alkyl or hetero (C_{8-10}) bicycloaryl (C_{0-6}) alkyl;
- 10 R^1 is hydrogen or (C_{1-6}) alkyl and R^2 is selected from a group consisting of hydrogen,
 cyano, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$, $-X^5NR^{12}C(O)OR^{12}$, $-R^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$,
 $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$, $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$,
 $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$, $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$,
 $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$, $-X^5S(O)_2R^{13}$, $-R^{14}$, $-X^5OR^{14}$, $-X^5SR^{14}$,
 15 $-X^5S(O)R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$, $-X^5OC(O)R^{14}$, $-X^5NR^{14}R^{12}$,
 $-X^5NR^{12}C(O)R^{14}$, $-X^5NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{14}R^{12}$, $-X^5NR^{12}S(O)_2R^{14}$,
 $-X^5NR^{12}C(O)NR^{14}R^{12}$ and $-X^5NR^{12}C(NR^{12})NR^{14}R^{12}$, wherein X^5 , R^{12} , R^{13} and R^{14} are as
 defined above; or R^1 and R^2 taken together with the carbon atom to which both R^1 and R^2 are
 attached form (C_{3-8}) cycloalkylene or (C_{3-8}) heterocycloalkylene; wherein within said R^2 any
 20 heteroaryl, aryl, cycloalkyl, heterocycloalkyl, cycloalkylene or heterocycloalkylene is
 unsubstituted or substituted with 1 to 3 radicals independently selected from (C_{1-6}) alkyl,
 (C_{1-6}) alkylidene, cyano, halo, halo-substituted (C_{1-4}) alkyl, nitro, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$,
 $-X^5NR^{12}C(O)OR^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$,
 $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$,
 25 $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$,
 $-X^5S(O)_2R^{13}$ and $-X^5C(O)R^{13}$, wherein X^5 , R^{12} and R^{13} are as defined above;
- R^3 is (C_{1-6}) alkyl or $-C(R^6)(R^6)X^6$, wherein R^6 is hydrogen or (C_{1-6}) alkyl and X^6 is
 selected from $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$, $-X^5NR^{12}C(O)OR^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$,
 $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$, $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$,
 30 $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$, $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$,
 $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5C(O)R^{13}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$, $-X^5S(O)_2R^{13}$, $-R^{14}$,
 $-X^5OR^{14}$, $-X^5SR^{14}$, $-X^5S(O)R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$, $-X^5OC(O)R^{14}$,
 $-X^5NR^{14}R^{12}$, $-X^5NR^{12}C(O)R^{14}$, $-X^5NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{14}R^{12}$, $-X^5S(O)_2NR^{14}R^{12}$,

-165-

$-X^5NR^{12}S(O)_2R^{14}$, $-X^5NR^{12}C(O)NR^{14}R^{12}$ and $-X^5NR^{12}C(NR^{12})NR^{14}R^{12}$ wherein X^5 , R^{12} , R^{13} and R^{14} are as defined above;

R^4 is selected from $-X^8NR^{12}R^{12}$, $-X^8NR^{12}C(O)R^{12}$, $-X^8NR^{12}C(O)OR^{12}$,
 $-X^8NR^{12}C(O)NR^{12}R^{12}$, $-X^8NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^8OR^{12}$, $-X^8SR^{12}$, $-X^5C(O)OR^{12}$,
 5 $-X^5C(O)R^{12}$, $-X^8OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^8S(O)_2NR^{12}R^{12}$, $-X^8NR^{12}S(O)_2R^{12}$,
 $-X^8P(O)(OR^{12})OR^{12}$, $-X^8OP(O)(OR^{12})OR^{12}$, $-X^5C(O)R^{13}$, $-X^8NR^{12}C(O)R^{13}$, $-X^8S(O)R^{13}$,
 $-X^8S(O)_2R^{13}$, $-R^{14}$, $-X^8OR^{14}$, $-X^8SR^{14}$, $-X^8S(O)R^{14}$, $-X^8S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$,
 $-X^8OC(O)R^{14}$, $-X^8NR^{14}R^{12}$, $-X^8NR^{12}C(O)R^{14}$, $-X^8NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{14}R^{12}$,
 $-X^8S(O)_2NR^{14}R^{12}$, $-X^8NR^{12}S(O)_2R^{14}$, $-X^8NR^{12}C(O)NR^{14}R^{12}$ and $-X^8NR^{12}C(NR^{12})NR^{14}R^{12}$
 10 wherein X^8 is (C_{1-6}) alkylene and X^5 , R^{12} , R^{13} and R^{14} are as defined above, with the proviso
 that when X^3 is cyano and X^2 is $-OR^4$, where R^4 is defined as $-R^{14}$, then R^{14} is
 (C_{3-10}) cycloalkyl (C_{1-6}) alkyl, hetero (C_{3-10}) cycloalkyl (C_{1-3}) alkyl, (C_{6-10}) aryl (C_{1-6}) alkyl,
 hetero (C_{5-10}) aryl (C_{1-6}) alkyl, (C_{9-10}) bicycloaryl (C_{1-6}) alkyl or
 hetero (C_{8-10}) bicycloaryl (C_{1-6}) alkyl;

15 R^{15} is (C_{6-10}) aryl, hetero (C_{5-10}) aryl, (C_{9-10}) bicycloaryl or hetero (C_{8-10}) bicycloaryl;

R^{17} is (C_{1-6}) alkyl, (C_{3-10}) cycloalkyl (C_{1-6}) alkyl, hetero (C_{3-10}) cycloalkyl (C_{1-6}) alkyl,
 (C_{6-10}) aryl (C_{1-6}) alkyl, hetero (C_{5-10}) aryl (C_{1-6}) alkyl, (C_{9-10}) bicycloaryl (C_{1-6}) alkyl or
 hetero (C_{8-10}) bicycloaryl (C_{1-6}) alkyl;

R^{18} is (C_{1-6}) alkyl, (C_{3-10}) cycloalkyl (C_{1-6}) alkyl, hetero (C_{3-10}) cycloalkyl (C_{1-6}) alkyl,
 20 (C_{6-10}) aryl (C_{1-6}) alkyl, hetero (C_{5-10}) aryl (C_{1-6}) alkyl, (C_{9-10}) bicycloaryl (C_{1-6}) alkyl or
 hetero (C_{8-10}) bicycloaryl (C_{1-6}) alkyl; and

R^{19} and R^{20} together with the atoms to which R^{19} and R^{20} are attached form
 (C_{4-8}) heterocycloalkylene, wherein no more than one of the ring member atoms comprising
 the ring is a heteroatom selected from $-NR^{21}-$ or $-O-$, wherein the ring is unsubstituted or
 25 substituted with R^2 , wherein R^2 is as defined above, and R^{21} is hydrogen, $-C(O)OR^{12}$,
 $-C(O)R^{12}$, $-C(O)NR^{12}R^{12}$, $-S(O)_2NR^{12}R^{12}$, $-S(O)R^{13}$ and $-S(O)_2R^{13}$, $-S(O)R^{14}$, $-S(O)_2R^{14}$,
 $-C(O)R^{14}$, $-C(O)OR^{14}$, $-C(O)NR^{12}R^{12}$ and $-S(O)_2NR^{14}R^{12}$, wherein R^{12} , R^{13} and R^{14} are as
 defined above;

wherein within R^3 , R^4 , R^{15} , R^{17} and R^{18} any alicyclic or aromatic ring system is
 30 unsubstituted or substituted further by 1 to 5 radicals independently selected from (C_{1-6}) alkyl,
 (C_{1-6}) alkylidene, cyano, halo, halo-substituted (C_{1-4}) alkyl, nitro, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$,
 $-X^5NR^{12}C(O)OR^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$,
 $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$,

-166-

$-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$,
 $-X^5C(O)R^{13}$ and $-X^5S(O)_2R^{13}$ and/or 1 radical selected from $-R^{14}$, $-X^5OR^{14}$, $-X^5SR^{14}$,
 $-X^5S(O)R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$, $-X^5OC(O)R^{14}$, $-X^5NR^{14}R^{12}$,
 $-X^5NR^{12}C(O)R^{14}$, $-X^5NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{14}R^{12}$, $-X^5S(O)_2NR^{14}R^{12}$, $-X^5NR^{12}S(O)_2R^{14}$,
5 $-X^5NR^{12}C(O)NR^{14}R^{12}$ and $-X^5NR^{12}C(NR^{12})NR^{14}R^{12}$; and within R^3 and R^4 any aliphatic
moiety is unsubstituted or substituted further by 1 to 5 radicals independently selected from
cyano, halo, nitro, $-NR^{12}R^{12}$, $-NR^{12}C(O)R^{12}$, $-NR^{12}C(O)OR^{12}$, $-NR^{12}C(O)NR^{12}R^{12}$,
 $-NR^{12}C(NR^{12})NR^{12}R^{12}$, $-OR^{12}$, $-SR^{12}$, $-C(O)OR^{12}$, $-C(O)R^{12}$, $-OC(O)R^{12}$, $-C(O)NR^{12}R^{12}$,
 $-S(O)_2NR^{12}R^{12}$, $-NR^{12}S(O)_2R^{12}$, $-P(O)(OR^{12})OR^{12}$, $-OP(O)(OR^{12})OR^{12}$, $-NR^{12}C(O)R^{13}$,
10 $-S(O)R^{13}$ and $-S(O)_2R^{13}$; wherein X^5 , R^{12} , R^{13} and R^{14} are as described above, with the proviso
that when X^2 is $-OR^4$, where R^4 is defined as $-R^{14}$, or $-NHR^{18}$, then any aromatic ring system
present within R^{14} or R^{18} is not substituted further by halo, (C_{3-10}) cycloalkyl,
hetero (C_{3-10}) cycloalkyl, (C_{6-10}) aryl, hetero (C_{5-10}) aryl, (C_{9-10}) bicycloaryl or
hetero (C_{8-10}) bicycloaryl; with the proviso that only one bicyclic ring structure is present
15 within R^3 , R^4 or R^{15} ; and the *N*-oxide derivatives, prodrug derivatives, protected derivatives,
individual isomers and mixtures of isomers thereof; and the pharmaceutically acceptable salts
and solvates of such compounds and the *N*-oxide derivatives, prodrug derivatives, protected
derivatives, individual isomers and mixtures of isomers thereof.

20 6. A compound of Claim 1 or 2 in which:
 X^1 is $-NHC(R^1)(R^2)X^3$ or $-NHCH(R^{19})C(O)R^{20}$;
 X^2 is $-OH$, $-OC(O)NR^{12}R^{12}$ or $-OC(O)R^{14}$, wherein R^{12} and R^{14} are as defined below;
 X^3 is cyano, $-C(R^7)(R^8)R^{16}$, $-C(R^6)(OR^6)_2$, $-CH_2C(O)R^{16}$, $-CH=CHS(O)_2R^5$,
 $-C(O)CF_2C(O)NR^5R^5$, $-C(O)C(O)NR^5R^6$, $-C(O)C(O)OR^5$, $-C(O)CH_2OR^5$,
25 $-C(O)CH_2N(R^6)SO_2R^5$ or $-C(O)C(O)R^5$; wherein R^5 is hydrogen, (C_{1-4}) alkyl,
 (C_{3-10}) cycloalkyl (C_{0-6}) alkyl, hetero (C_{3-10}) cycloalkyl (C_{0-3}) alkyl, (C_{6-10}) aryl (C_{0-6}) alkyl,
hetero (C_{5-10}) aryl (C_{0-6}) alkyl, (C_{9-10}) bicycloaryl (C_{0-6}) alkyl or
hetero (C_{8-10}) bicycloaryl (C_{0-6}) alkyl; R^6 is hydrogen, hydroxy or (C_{1-6}) alkyl; or where X^3
contains an $-NR^5R^6$ group, R^5 and R^6 together with the nitrogen atom to which they are both
30 attached, form hetero (C_{3-10}) cycloalkyl, hetero (C_{5-10}) aryl or hetero (C_{8-10}) bicycloaryl; R^7 is
hydrogen or (C_{1-4}) alkyl and R^8 is hydroxy or R^7 and R^8 together form oxo; R^{16} is hydrogen, -
 X^4 , $-CF_3$, $-CF_2CF_2R^9$ or $-N(R^6)OR^6$; R^9 is hydrogen, halo, (C_{1-4}) alkyl, (C_{5-10}) aryl (C_{0-6}) alkyl or
 (C_{5-10}) heteroaryl (C_{0-6}) alkyl;

-167-

X^4 comprises a heteromonocyclic ring containing 4 to 7 ring member atoms or a fused heterobicyclic ring system containing 8 to 14 ring member atoms and any carbocyclic ketone, iminoketone or thioketone derivative thereof;

wherein within R^5 , X^3 or X^4 any alicyclic or aromatic ring system is unsubstituted or substituted further by 1 to 5 radicals independently selected from (C₁₋₆)alkyl, (C₁₋₆)alkylidene, cyano, halo, halo-substituted(C₁₋₄)alkyl, nitro, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$, $-X^5NR^{12}C(O)OR^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$, $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$, $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$ and $-X^5S(O)_2R^{13}$ and/or 1 radical selected from $-R^{14}$, $-X^5OR^{14}$, $-X^5SR^{14}$, $-X^5S(O)R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$, $-X^5OC(O)R^{14}$, $-X^5NR^{14}R^{12}$, $-X^5NR^{12}C(O)R^{14}$, $-X^5NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{14}R^{12}$, $-X^5NR^{12}S(O)_2R^{14}$, $-X^5NR^{12}C(O)NR^{14}R^{12}$ and $-X^5NR^{12}C(NR^{12})NR^{14}R^{12}$, wherein X^5 is a bond or (C₁₋₆)alkylene; R^{12} at each occurrence independently is hydrogen, (C₁₋₆)alkyl or halo-substituted(C₁₋₆)alkyl; R^{13} is (C₁₋₆)alkyl or halo-substituted(C₁₋₆)alkyl; and R^{14} is (C₃₋₁₀)cycloalkyl(C₀₋₆)alkyl, hetero(C₃₋₁₀)cycloalkyl(C₀₋₃)alkyl, (C₆₋₁₀)aryl(C₀₋₆)alkyl, hetero(C₅₋₁₀)aryl(C₀₋₆)alkyl, (C₉₋₁₀)bicycloaryl(C₀₋₆)alkyl or hetero(C₈₋₁₀)bicycloaryl(C₀₋₆)alkyl;

R^1 is hydrogen or (C₁₋₆)alkyl and R^2 is selected from a group consisting of hydrogen, cyano, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$, $-X^5NR^{12}C(O)OR^{12}$, $-R^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$, $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$, $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$, $-X^5S(O)_2R^{13}$, $-R^{14}$, $-X^5OR^{14}$, $-X^5SR^{14}$, $-X^5S(O)R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$, $-X^5OC(O)R^{14}$, $-X^5NR^{14}R^{12}$, $-X^5NR^{12}C(O)R^{14}$, $-X^5NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{14}R^{12}$, $-X^5NR^{12}S(O)_2R^{14}$, $-X^5NR^{12}C(O)NR^{14}R^{12}$ and $-X^5NR^{12}C(NR^{12})NR^{14}R^{12}$, wherein X^5 , R^{12} , R^{13} and R^{14} are as defined above; or R^1 and R^2 taken together with the carbon atom to which both R^1 and R^2 are attached form (C₃₋₈)cycloalkylene or (C₃₋₈)heterocycloalkylene; wherein within said R^2 any heteroaryl, aryl, cycloalkyl, heterocycloalkyl, cycloalkylene or heterocycloalkylene is unsubstituted or substituted with 1 to 3 radicals independently selected from (C₁₋₆)alkyl, (C₁₋₆)alkylidene, cyano, halo, halo-substituted(C₁₋₄)alkyl, nitro, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$, $-X^5NR^{12}C(O)OR^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$, $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$, $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$,

-168-

$-X^5S(O)_2R^{13}$ and $-X^5C(O)R^{13}$, wherein X^5 , R^{12} and R^{13} are as defined above;

R^3 is (C_{1-6}) alkyl or $-C(R^6)(R^6)X^6$, wherein R^6 is hydrogen or (C_{1-6}) alkyl and X^6 is selected from $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$, $-X^5NR^{12}C(O)OR^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$, $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$, $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5C(O)R^{13}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$, $-X^5S(O)_2R^{13}$, $-R^{14}$, $-X^5OR^{14}$, $-X^5SR^{14}$, $-X^5S(O)R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$, $-X^5OC(O)R^{14}$, $-X^5NR^{14}R^{12}$, $-X^5NR^{12}C(O)R^{14}$, $-X^5NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{14}R^{12}$, $-X^5S(O)_2NR^{14}R^{12}$, $-X^5NR^{12}S(O)_2R^{14}$, $-X^5NR^{12}C(O)NR^{14}R^{12}$ and $-X^5NR^{12}C(NR^{12})NR^{14}R^{12}$ wherein X^5 , R^{12} , R^{13} and R^{14} are as defined above; and

R^{19} and R^{20} together with the atoms to which R^{19} and R^{20} are attached form (C_{4-8}) heterocycloalkylene, wherein no more than one of the ring member atoms comprising the ring is a heteroatom selected from $-NR^{21}-$ or $-O-$, wherein the ring is unsubstituted or substituted with R^2 , wherein R^2 is as defined above, and R^{21} is hydrogen, $-C(O)OR^{12}$, $-C(O)R^{12}$, $-C(O)NR^{12}R^{12}$, $-S(O)_2NR^{12}R^{12}$, $-S(O)R^{13}$ and $-S(O)_2R^{13}$, $-S(O)R^{14}$, $-S(O)_2R^{14}$, $-C(O)R^{14}$, $-C(O)OR^{14}$, $-C(O)NR^{12}R^{12}$ and $-S(O)_2NR^{14}R^{12}$, wherein R^{12} , R^{13} and R^{14} are as defined above;

wherein within R^3 , R^4 , R^{15} , R^{17} and R^{18} any alicyclic or aromatic ring system is unsubstituted or substituted further by 1 to 5 radicals independently selected from (C_{1-6}) alkyl, (C_{1-6}) alkylidene, cyano, halo, halo-substituted (C_{1-4}) alkyl, nitro, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$, $-X^5NR^{12}C(O)OR^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$, $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$, $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$, $-X^5C(O)R^{13}$ and $-X^5S(O)_2R^{13}$ and/or 1 radical selected from $-R^{14}$, $-X^5OR^{14}$, $-X^5SR^{14}$, $-X^5S(O)R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$, $-X^5OC(O)R^{14}$, $-X^5NR^{14}R^{12}$, $-X^5NR^{12}C(O)R^{14}$, $-X^5NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{14}R^{12}$, $-X^5S(O)_2NR^{14}R^{12}$, $-X^5NR^{12}S(O)_2R^{14}$, $-X^5NR^{12}C(O)NR^{14}R^{12}$ and $-X^5NR^{12}C(NR^{12})NR^{14}R^{12}$; and within R^3 and R^4 any aliphatic moiety is unsubstituted or substituted further by 1 to 5 radicals independently selected from cyano, halo, nitro, $-NR^{12}R^{12}$, $-NR^{12}C(O)R^{12}$, $-NR^{12}C(O)OR^{12}$, $-NR^{12}C(O)NR^{12}R^{12}$, $-NR^{12}C(NR^{12})NR^{12}R^{12}$, $-OR^{12}$, $-SR^{12}$, $-C(O)OR^{12}$, $-C(O)R^{12}$, $-OC(O)R^{12}$, $-C(O)NR^{12}R^{12}$, $-S(O)_2NR^{12}R^{12}$, $-NR^{12}S(O)_2R^{12}$, $-P(O)(OR^{12})OR^{12}$, $-OP(O)(OR^{12})OR^{12}$, $-NR^{12}C(O)R^{13}$, $-S(O)R^{13}$ and $-S(O)_2R^{13}$; wherein X^5 , R^{12} , R^{13} and R^{14} are as described above; with the proviso that only one bicyclic ring structure is present within R^3 , R^4 or R^{15} ; and the *N*-oxide

-169-

derivatives, prodrug derivatives, protected derivatives, individual isomers and mixtures of isomers thereof; and the pharmaceutically acceptable salts and solvates of such compounds and the *N*-oxide derivatives, prodrug derivatives, protected derivatives, individual isomers and mixtures of isomers thereof.

5

7. The compound of Claim 1 or Claim 2 in which:

X^1 is $-NHC(R^1)(R^2)C(O)C(O)NR^5R^6$, wherein R^5 is hydrogen, (C_{1-4}) alkyl, (C_{3-10}) cycloalkyl (C_{0-6}) alkyl, hetero (C_{3-10}) cycloalkyl (C_{0-3}) alkyl, (C_{6-10}) aryl (C_{0-6}) alkyl, hetero (C_{5-10}) aryl (C_{0-6}) alkyl, (C_{9-10}) bicycloaryl (C_{0-6}) alkyl or
 10 hetero (C_{8-10}) bicycloaryl (C_{0-6}) alkyl and R^6 is hydrogen, hydroxy or (C_{1-6}) alkyl or R^5 and R^6 together with the nitrogen atom to which they are both attached form hetero (C_{3-10}) cycloalkyl, hetero (C_{5-10}) aryl or hetero (C_{8-10}) bicycloaryl;

X^2 is hydrogen;

wherein within X^1 any alicyclic or aromatic ring system is unsubstituted or substituted
 15 further by 1 to 5 radicals independently selected from (C_{1-6}) alkyl, (C_{1-6}) alkylidene, cyano, halo, halo-substituted (C_{1-4}) alkyl, nitro, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$, $-X^5NR^{12}C(O)OR^{12}$, $-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$, $-X^5C(O)OR^{12}$, $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$, $-X^5NR^{12}S(O)_2R^{12}$, $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$ and $-X^5S(O)_2R^{13}$
 20 and/or 1 radical selected from $-R^{14}$, $-X^5OR^{14}$, $-X^5SR^{14}$, $-X^5S(O)R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)OR^{14}$, $-X^5OC(O)R^{14}$, $-X^5NR^{14}R^{12}$, $-X^5NR^{12}C(O)R^{14}$, $-X^5NR^{12}C(O)OR^{14}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{14}R^{12}$, $-X^5NR^{12}S(O)_2R^{14}$, $-X^5NR^{12}C(O)NR^{14}R^{12}$ and $-X^5NR^{12}C(NR^{12})NR^{14}R^{12}$, wherein X^5 is a bond or (C_{1-6}) alkylene; R^{12} at each occurrence independently is hydrogen, (C_{1-6}) alkyl or halo-substituted (C_{1-6}) alkyl; R^{13} is (C_{1-6}) alkyl or
 25 halo-substituted (C_{1-6}) alkyl; and R^{14} is (C_{3-10}) cycloalkyl (C_{0-6}) alkyl, hetero (C_{3-10}) cycloalkyl (C_{0-3}) alkyl, (C_{6-10}) aryl (C_{0-6}) alkyl, hetero (C_{5-10}) aryl (C_{0-6}) alkyl, (C_{9-10}) bicycloaryl (C_{0-6}) alkyl or hetero (C_{8-10}) bicycloaryl (C_{0-6}) alkyl;

R^1 is hydrogen and R^2 is (C_{1-6}) alkyl; and

R^3 is $-CH_2X^6$, wherein X^6 is $-X^5NR^{12}S(O)_2R^{12}$ or $-X^5S(O)_2R^{14}$ wherein X^5 , R^{12} and R^{14}
 30 are as defined above;

wherein within R^3 any alicyclic or aromatic ring system is unsubstituted or substituted further by 1 to 5 radicals independently selected from (C_{1-6}) alkyl, (C_{1-6}) alkylidene, cyano, halo, halo-substituted (C_{1-4}) alkyl, nitro, $-X^5NR^{12}R^{12}$, $-X^5NR^{12}C(O)R^{12}$, $-X^5NR^{12}C(O)OR^{12}$,

-170-

$-X^5NR^{12}C(O)NR^{12}R^{12}$, $-X^5NR^{12}C(NR^{12})NR^{12}R^{12}$, $-X^5OR^{12}$, $-X^5SR^{12}$, $-X^5C(O)OR^{12}$,
 $-X^5C(O)R^{12}$, $-X^5OC(O)R^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2NR^{12}R^{12}$, $-X^5NR^{12}S(O)_2R^{12}$,
 $-X^5P(O)(OR^{12})OR^{12}$, $-X^5OP(O)(OR^{12})OR^{12}$, $-X^5NR^{12}C(O)R^{13}$, $-X^5S(O)R^{13}$, $-X^5C(O)R^{13}$ and
 $-X^5S(O)_2R^{13}$ and within R^3 any aliphatic moiety is unsubstituted or substituted further by 1 to
5 5 radicals independently selected from cyano, halo, nitro, $-NR^{12}R^{12}$, $-NR^{12}C(O)R^{12}$,
 $-NR^{12}C(O)OR^{12}$, $-NR^{12}C(O)NR^{12}R^{12}$, $-NR^{12}C(NR^{12})NR^{12}R^{12}$, $-OR^{12}$, $-SR^{12}$, $-C(O)OR^{12}$,
 $-C(O)R^{12}$, $-OC(O)R^{12}$, $-C(O)NR^{12}R^{12}$, $-S(O)_2NR^{12}R^{12}$, $-NR^{12}S(O)_2R^{12}$, $-P(O)(OR^{12})OR^{12}$,
 $-OP(O)(OR^{12})OR^{12}$, $-NR^{12}C(O)R^{13}$, $-S(O)R^{13}$ and $-S(O)_2R^{13}$; wherein X^5 , R^{12} , R^{13} and R^{14} are
as described above; with the proviso that only one bicyclic ring structure is present within R^3 ;
10 and the *N*-oxide derivatives, prodrug derivatives, protected derivatives, individual isomers and
mixtures of isomers thereof; and the pharmaceutically acceptable salts and solvates of such
compounds and the *N*-oxide derivatives, prodrug derivatives, protected derivatives, individual
isomers and mixtures of isomers thereof.

15 8. The compound of Claim 3 in which:

X^1 is $-NHC(R^1)(R^2)X^3$ or $-NHCH(R^{19})C(O)R^{20}$, wherein R^1 is hydrogen or
 (C_{1-6}) alkyl and R^2 is hydrogen, (C_{1-6}) alkyl, $-X^5OR^{12}$, $-X^5S(O)R^{13}$, $-X^5OR^{14}$,
 (C_{6-10}) aryl (C_{0-6}) alkyl or hetero (C_{5-10}) aryl (C_{0-6}) alkyl or R^1 and R^2 taken together with
the carbon atom to which both R^1 and R^2 are attached form (C_{3-6}) cycloalkylene or
20 (C_{3-6}) heterocycloalkylene, wherein within said R^2 any heteroaryl, aryl, cycloalkylene
or heterocycloalkylene is unsubstituted or substituted with (C_{1-6}) alkyl or hydroxy,
wherein X^3 is cyano, $-C(O)R^{16}$, $-C(R^6)(OR^6)_2$, $-CH=CHS(O)_2R^5$, $-CH_2C(O)R^{16}$,
 $-C(O)CF_2C(O)NR^5R^5$, $-C(O)C(O)NR^5R^6$, $-C(O)C(O)OR^5$, $-C(O)CH_2OR^5$,
 $-C(O)CH_2N(R^6)SO_2R^5$ or $-C(O)C(O)R^5$ and R^{19} and R^{20} together with the atoms to
25 which R^{19} and R^{20} are attached form (C_{4-8}) heterocycloalkylene, wherein no more than
one of the ring member atoms comprising the ring is a heteroatom selected from
 $-NR^{21}$ or $-O-$, wherein the ring is unsubstituted or substituted with (C_{1-6}) alkyl or
 $-X^5C(O)OR^{12}$ and R^{21} is hydrogen, (C_{1-6}) alkyl, $-X^5C(O)R^{12}$, $-X^5C(O)OR^{12}$, $-R^{14}$,
 $-X^5C(O)R^{14}$ or $-C(O)OR^{14}$;

30 X^2 is $-OH$ or $-OC(O)NR^{12}R^{12}$, wherein each R^{12} independently represent
hydrogen or (C_{1-6}) alkyl, wherein said alkyl is unsubstituted or substituted with
hydroxy or methoxy, or X^2 is $-OC(O)NHR^{14}$, wherein R^{14} is
 (C_{3-10}) cycloalkyl (C_{0-6}) alkyl or hetero (C_{3-10}) cycloalkyl (C_{1-3}) alkyl, or X^2 is $-OC(O)R^{14}$,

-171-

wherein R^{14} is $-NR^{22}R^{23}$ and R^{22} and R^{23} together with the nitrogen atom to which both R^{22} and R^{23} attached form a hetero(C_{4-6})cycloalkyl ring, which ring may be unsubstituted or substituted with hydroxy; and

R^3 is $-CH_2X^6$; wherein X^6 is selected from $-X^5SR^{12}$, $-X^5C(O)NR^{12}R^{12}$, $-X^5S(O)_2R^{13}$, $-X^5C(O)R^{13}$, $-X^5OR^{12}$, $-X^5SR^{14}$, $-X^5R^{14}$, $-X^5S(O)_2R^{14}$, $-X^5C(O)R^{14}$, $-X^5C(O)NR^{14}R^{12}$; and the *N*-oxide derivatives, prodrug derivatives, protected derivatives, individual isomers and mixtures of isomers thereof; and the pharmaceutically acceptable salts and solvates of such compounds and the *N*-oxide derivatives, prodrug derivatives, protected derivatives, individual isomers and mixtures of isomers thereof.

9. The compound of Claim 8 in which:

X^3 is cyano, $-C(O)X^4$, $-C(O)H$, $-C(O)N(CH_3)OCH_3$, $-CH(OCH_3)_2$, $-C(O)CF_3$, $-C(O)CF_2CF_3$, $-CH_2C(O)R^{16}$, (E)-2-benzenesulfonyl-vinyl, 2-dimethylcarbamoyl-2,2-difluoro-acetyl, 2-oxo-2-pyrrolidin-1-yl-acetyl, 2-morpholin-4-yl-2-oxo-acetyl, 2-oxo-2-piperazin-1-yl-acetyl, 2-(4-methanesulfonylpiperazin-1-yl)-2-oxo-acetyl, 2-(1,1-dioxo-1 λ^6 -thiomorpholin-4-yl)-2-oxo-acetyl, dimethylaminooxalyl, tetrahydro-pyran-4-ylaminooxalyl, 2-morpholin-4-ylethylaminooxalyl, cyclopentyl-ethyl-aminooxalyl, pyridin-3-ylaminooxalyl, phenylaminooxalyl, 1-benzoyl-piperidin-4-ylaminooxalyl, 1-benzylcarbamoyl-methanoyl, 1-benzyloxy(oxalyl), 2-benzyloxy-acetyl, 2-benzenesulfonylamino-ethanoyl, 2-oxo-2-phenyl-ethanoyl, 3*H*-oxazole-2-carbonyl, 5-trifluoromethyl-oxazole-2-carbonyl, 3-trifluoromethyl-[1,2,4]oxadiazole-5-carbonyl, 2,2,3,3,3-pentafluoro-propionyl, hydroxyaminooxalyl, oxalyl, 2-(1,3-dihydro-isoindol-2-yl)-2-oxo-acetyl, benzothiazol-2-ylaminooxalyl, 2-oxo-ethyl, 2-oxazol-2-yl-2-oxo-ethyl or 2-benzooxazol-2-yl-2-oxo-ethyl;

X^2 is selected from $-OH$, dimethylcarbamoyloxy, morpholin-4-ylcarbonyloxy, piperidin-1-yl-carbonyloxy, pyrrolidin-1-yl-carbonyloxy, pyrimidin-2-ylamino, tetrahydro-pyran-4-ylamino, 1-methyl-piperidin-4-ylamino, *N*-(2-methoxyethyl)-*N*-(tetrahydro-pyran-4-yl)amino, isopropylamino and cyclohexylamino; 4-*tert*-butoxycarbonylpiperazin-1-ylcarbonyloxy, *N*-benzyl-carbamoyloxy, pyrrolidin-1-yl-carbonyloxy, *N,N*-dimethyl-carbamoyloxy, piperidin-1-yl-carbonyloxy, 4-methanesulfonylpiperazin-1-yl-carbonyloxy, 4-ethoxycarbonylpiperazin-1-

-172-

ylcarbonyloxy, *N*-cyclohexyl-carbamoyloxy, *N*-phenyl-carbamoyloxy, *N*-(5,6,7,8-tetrahydro-naphthalen-1-yl)-carbamoyloxy, *N*-butyl-*N*-methyl-carbamoyloxy, *N*-pyridin-3-yl-carbamoyloxy, *N*-isopropyl-carbamoyloxy, *N*-pyridin-4-yl-carbamoyloxy, *N*-cyanomethyl-*N*-methyl-carbamoyloxy, *N,N*-bis-(2-methoxy-ethyl)-carbamoyloxy, *N*-phenethyl-carbamoyloxy, piperazine-carbonyloxy, *N*-naphthalen-2-yl-carbamoyloxy, 4-benzyl-piperazine-1-carbamoyloxy, 4-(1-furan-2-yl-carbonyl)-piperazine-1-carbamoyloxy, thiomorpholin-4-yl-carbonyloxy, 1,1-dioxo-1λ⁶-thiomorpholin-4-yl-carbonyloxy, bis-(2-methoxy-ethyl)-carbamoyloxy, morpholin-4-ylcarbonyloxy, 2-methoxyethylcarbamoyloxy, diethylcarbamoyloxy, pyrrolidin-1-ylcarbonyloxy, 2-hydroxyethylcarbamoyloxy, tetrahydro-furan-2-ylmethylcarbamoyloxy, cyclopropylcarbamoyloxy, *tert*-butylcarbamoyloxy, 3-hydroxy-pyrrolidin-1-yl-carbonyloxy and carbamoyloxy; and

R^3 is thiophene-2-sulfonyl-methyl, 3-chloro-2-fluoro-phenyl-methane-sulfonyl-methyl, benzene-sulfonyl-methyl, phenyl-methane-sulfonyl-methyl,

2-(1,1-difluoro-methoxy)-phenyl-methane-sulfonyl-methyl, 2-benzene-sulfonyl-ethyl, 2-(pyridine-2-sulfonyl)-ethyl, 2-(pyridine-4-sulfonyl)-ethyl, 2-phenyl-methanesulfonyl-ethyl, oxy-pyridin-2-yl-methane-sulfonyl-methyl, prop-2-ene-1-sulfonyl-methyl, 4-methoxy-phenyl-methane-sulfonyl-methyl, *p*-tolyl-methane-sulfonyl-methyl, 4-chloro-phenyl-methane-sulfonyl-methyl, *o*-tolyl-methane-sulfonyl-methyl, 3,5-dimethyl-phenyl-methane-sulfonyl-methyl, 4-trifluoro-methyl-phenyl-methane-sulfonyl-methyl, 4-trifluoro-methoxy-phenyl-methane-sulfonyl-methyl, 2-bromo-phenyl-methane-sulfonyl-methyl, pyridin-2-yl-methane-sulfonyl-methyl, pyridin-3-yl-methane-sulfonyl-methyl, pyridin-4-yl-methane-sulfonyl-methyl, naphthalen-2-yl-methane-sulfonyl-methyl, 3-methyl-phenyl-methane-sulfonyl-methyl, 3-trifluoro-methyl-phenyl-methane-sulfonyl-methyl, 3-trifluoro-methoxy-phenyl-methane-sulfonyl-methyl, 4-fluoro-2-trifluoromethoxy-phenyl-methane-sulfonylmethyl, 2-fluoro-6-trifluoromethyl-phenylmethanesulfonylmethyl, 3-chloro-phenylmethanesulfonylmethyl, 2-fluoro-phenylmethanesulfonylmethyl, 2-trifluoro-phenylmethanesulfonylmethyl, 2-cyano-phenylmethanesulfonylmethyl, 4-*tert*-butyl-phenylmethanesulfonylmethyl, 2-fluoro-3-methyl-phenyl-methane-sulfonyl-methyl, 3-fluoro-phenylmethanesulfonylmethyl, 4-fluoro-phenylmethane-sulfonylmethyl, 2-chloro-phenylmethanesulfonylmethyl, 2,5-difluoro-phenylmethane-sulfonylmethyl, 2,6-difluoro-phenylmethanesulfonylmethyl, 2,5-dichloro-phenyl-

-173-

methane-sulfonylmethyl, 3,4-dichloro-phenylmethanesulfonylmethyl,
2-(1,1-difluoro-methoxy)-phenyl-methanesulfonylmethyl, 2-cyano-phenyl-methane-
sulfonyl-methyl, 3-cyano-phenylmethanesulfonylmethyl, 2-trifluoro-methoxy-phenyl-
methane-sulfonylmethyl, 2,3-difluoro-phenylmethanesulfonylmethyl,
5 2,5-difluoro-phenyl-methanesulfonylmethyl, biphenyl-2-ylmethanesulfonylmethyl,
cyclohexylmethyl, 3-fluoro-phenyl-methanesulfonylmethyl, 3,4-difluoro-phenyl-
methanesulfonylmethyl, 2,4-difluoro-phenylmethanesulfonylmethyl, 2,4,6-trifluoro-
phenylmethanesulfonylmethyl, 2,4,5-trifluoro-phenylmethanesulfonylmethyl,
2,3,4-trifluoro-phenylmethanesulfonylmethyl, 2,3,5-trifluoro-phenyl-methane-
10 sulfonylmethyl, 2,5,6-trifluoro-phenylmethanesulfonylmethyl, 2-chloro-5-trifluoro-
methylphenylmethanesulfonylmethyl, 2-methyl-propane-1-sulfonyl,
2-fluoro-3-trifluoro-methylphenylmethanesulfonylmethyl, 2-fluoro-4-trifluoro-
methylphenylmethanesulfonylmethyl, 2-fluoro-5-trifluoro-methyl-phenyl-methane-
sulfonyl-methyl, 4-fluoro-3-trifluoro-methylphenylmethanesulfonylmethyl,
15 2-methoxy-phenyl-methanesulfonylmethyl,
3,5-bis-trifluoromethyl-phenylmethanesulfonylmethyl,
4-difluoromethoxy-phenylmethanesulfonylmethyl, 2-difluoro-methoxy-phenyl-
methanesulfonylmethyl, 3-difluoromethoxy-phenylmethanesulfonylmethyl,
2,6-dichloro-phenylmethanesulfonylmethyl, biphenyl-4-ylmethanesulfonylmethyl,
20 3,5-dimethyl-isoxazol-4-ylmethanesulfonylmethyl, 5-chloro-thien-2-yl-methane-
sulfonylmethyl, 2-[4-(1,1-difluoro-methoxy)-benzenesulfonyl]-ethyl,
2-[2-(1,1-difluoro-methoxy)-benzenesulfonyl]-ethyl, 2-[3-(1,1-difluoro-
methoxy)-benzenesulfonyl]-ethyl, 2-(4-trifluoromethoxy-benzenesulfonyl)-ethyl,
2-(3-trifluoromethoxy-benzenesulfonyl)-ethyl, 2-(2-trifluoro-methoxy-benzene-
25 sulfonyl)-ethyl, (cyanomethyl-methyl-carbamoyl)-methyl, biphenyl-3-ylmethyl,
2-oxo-2-pyrrolidin-1-yl-ethyl, 2-benzenesulfonyl-ethyl, isobutylsulfanylmethyl,
2-phenylsulfanyl-ethyl, cyclohexylmethanesulfonylmethyl, 2-cyclohexyl-
ethanesulfonyl, benzyl, naphthalen-2-yl, benzylsulfanylmethyl,
2-trifluoromethyl-benzylsulfanylmethyl, phenylsulfanyl-ethyl, cyclopropyl-
30 methanesulfonylmethyl, 5-bromo-thien-2-ylmethyl, 3-phenyl-propyl, 2,2-difluoro-
3-phenyl-propyl, 3,4,5-trimethoxy-phenylmethanesulfonylmethyl, 2,2-difluoro-
3-thien-2-yl-propyl, cyclohexylethyl, cyclohexylmethyl, *tert*-butylmethyl,
1-methylcyclohexylmethyl, 1-methylcyclopentylmethyl, 2,2-difluoro-3-phenylpropyl,

-174-

2,2-dimethyl-3-phenylpropyl, 1-benzylcyclopropylmethyl, $-X^5S(O)_2R^{13}$ and $-X^5S(O)_2R^{14}$, wherein R^{13} is alkyl and R^{14} is phenyl which phenyl is unsubstituted or substituted; and the *N*-oxide derivatives, prodrug derivatives, protected derivatives, individual isomers and mixtures of isomers thereof; and the pharmaceutically acceptable salts and solvates of such compounds and the *N*-oxide derivatives, prodrug derivatives, protected derivatives, individual isomers and mixtures of isomers thereof.

10. A compound of Claim 9 in which:

X^3 is 1*H*-benzoimidazol-2-ylcarbonyl, pyrimidin-2-ylcarbonyl, benzooxazol-2-ylcarbonyl, benzothiazol-2-ylcarbonyl, pyridazin-3-ylcarbonyl, 3-phenyl-[1,2,4]oxadiazol-5-ylcarbonyl or 3-ethyl-[1,2,4]oxadiazol-5-ylcarbonyl, 2-oxo-2-pyrrolidin-1-yl-acetyl, 2-morpholin-4-yl-2-oxo-acetyl, 2-oxo-2-piperazin-1-yl-acetyl, 2-(4-methanesulfonyl-piperazin-1-yl)-2-oxo-acetyl, 2-(1,1-dioxo-1λ⁶-thiomorpholin-4-yl)-2-oxo-acetyl, dimethylaminooxalyl, tetrahydropyran-4-ylaminooxalyl, 2-morpholin-4-yl-ethylaminooxalyl, cyclopentyl-ethylaminooxalyl, pyridin-3-ylaminooxalyl, phenylaminooxalyl or 1-benzoylpiperidin-4-ylaminooxalyl;

X^2 is selected from -OH, dimethylcarbamoyloxy, morpholin-4-ylcarbonyloxy, piperidin-1-yl-carbonyloxy, pyrrolidin-1-yl-carbonyloxy, pyrimidin-2-ylamino, tetrahydro-pyran-4-ylamino, 1-methyl-piperidin-4-ylamino, *N*-(2-methoxyethyl)-*N*-(tetrahydro-pyran-4-yl)amino, isopropylamino and cyclohexylamino;

R^3 is cyclohexylethyl, cyclohexylmethyl, *tert*-butylmethyl, 1-methylcyclohexylmethyl, 1-methylcyclopentylmethyl, 2,2-difluoro-3-phenylpropyl, 2,2-dimethyl-3-phenylpropyl, 1-benzylcyclopropylmethyl, $-X^5S(O)_2R^{13}$ or $-X^5S(O)_2R^{14}$, wherein R^{13} is alkyl and R^{14} is phenyl which phenyl is unsubstituted or substituted; and the pharmaceutically acceptable salts and solvates of such compounds and the *N*-oxide derivatives, prodrug derivatives, protected derivatives, individual isomers and mixtures of isomers thereof.

11. The compound of Claim 3 in which:

X^1 is $-NHC(R^1)(R^2)X^3$ or $-NHCH(R^{19})C(O)R^{20}$, wherein R^1 is hydrogen or (C₁₋₆)alkyl and R^2 is hydrogen, (C₁₋₆)alkyl, $-X^5OR^{12}$, $-X^5S(O)R^{13}$, $-X^5OR^{14}$, (C₆₋₁₀)aryl(C₀₋₆)alkyl or hetero(C₅₋₁₀)aryl(C₀₋₆)alkyl or R^1 and R^2 taken together with

-175-

the carbon atom to which both R¹ and R² are attached form (C₃₋₆)cycloalkylene or (C₃₋₆)heterocycloalkylene, wherein within said R² any heteroaryl, aryl, cycloalkylene or heterocycloalkylene is unsubstituted or substituted with (C₁₋₆)alkyl or hydroxy, wherein X³ is cyano, -C(O)R¹⁶, -C(R⁶)(OR⁶)₂, -CH=CHS(O)₂R⁵, -CH₂C(O)R¹⁶,
 5 -C(O)CF₂C(O)NR⁵R⁵, -C(O)C(O)NR⁵R⁶, -C(O)C(O)OR⁵, -C(O)CH₂OR⁵,
 -C(O)CH₂N(R⁶)SO₂R⁵ or -C(O)C(O)R⁵ and R¹⁹ and R²⁰ together with the atoms to which R¹⁹ and R²⁰ are attached form (C₄₋₈)heterocycloalkylene, wherein no more than one of the ring member atoms comprising the ring is a heteroatom selected from -NR²¹ - or -O-, wherein the ring is unsubstituted or substituted with (C₁₋₆)alkyl or
 10 -X⁵C(O)OR¹² and R²¹ is hydrogen, (C₁₋₆)alkyl, -X⁵C(O)R¹², -X⁵C(O)OR¹², -R¹⁴,
 -X⁵C(O)R¹⁴ or -C(O)OR¹⁴;

X² is -NHR¹⁵, wherein R¹⁵ is (C₆₋₁₀)aryl, hetero(C₅₋₁₀)aryl, (C₉₋₁₀)bicycloaryl or hetero(C₈₋₁₀)bicycloaryl, or -NR¹⁷R¹⁸, wherein R¹⁷ is hetero(C₃₋₁₀)cycloalkyl and R¹⁸ is hydrogen or R¹⁷ and R¹⁸ independently are (C₆₋₁₀)aryl(C₁₋₆)alkyl or
 15 hetero(C₅₋₁₀)aryl(C₁₋₆)alkyl, wherein within R¹⁵, R¹⁷ and R¹⁸ any alicyclic or aromatic ring system is unsubstituted or substituted further by 1 to 5 radicals independently selected from (C₁₋₆)alkyl, cyano, halo, nitro, halo-substituted(C₁₋₄)alkyl, -X⁵OR¹²,
 -X⁵C(O)OR¹², -X⁵C(O)R¹³, -X⁵C(O)NR¹²R¹², -X⁵NR¹²S(O)₂R¹² and/or 1 radical selected from -R¹⁴, -X⁵OR¹⁴ and -X⁵C(O)NR¹⁴R¹²; and

20 R³ is -CH₂X⁶; wherein X⁶ is selected from -X⁵SR¹², -X⁵C(O)NR¹²R¹²,
 -X⁵S(O)₂R¹³, -X⁵C(O)R¹³, -X⁵OR¹², -X⁵SR¹⁴, -X⁵R¹⁴, -X⁵S(O)₂R¹⁴, -X⁵C(O)R¹⁴,
 -X⁵C(O)NR¹⁴R¹²; and the *N*-oxide derivatives, prodrug derivatives, protected derivatives, individual isomers and mixtures of isomers thereof; and the
 25 pharmaceutically acceptable salts and solvates of such compounds and the *N*-oxide derivatives, prodrug derivatives, protected derivatives, individual isomers and mixtures of isomers thereof.

12. The compound of Claim 11 in which:

30 X³ is cyano, -C(O)X⁴, -C(O)H, -C(O)N(CH₃)OCH₃, -CH(OCH₃)₂, -C(O)CF₃,
 -C(O)CF₂CF₃, -CH₂C(O)R¹⁶, (E)-2-benzenesulfonyl-vinyl,
 2-dimethylcarbamoyl-2,2-difluoro-acetyl, 2-oxo-2-pyrrolidin-1-yl-acetyl, 2-morpholin-4-yl-2-oxo-acetyl, 2-oxo-2-piperazin-1-yl-acetyl, 2-(4-methanesulfonyl-piperazin-1-yl)-2-oxo-acetyl, 2-(1,1-dioxo-1λ⁶-thiomorpholin-4-yl)-2-oxo-acetyl,

-176-

dimethylaminooxalyl, tetrahydro-pyran-4-ylaminooxalyl, 2-morpholin-4-yl-ethylaminooxalyl, cyclopentyl-ethyl-aminooxalyl, pyridin-3-ylaminooxalyl, phenylaminooxalyl, 1-benzoyl-piperidin-4-ylaminooxalyl, 1-benzylcarbamoyl-methanoyl, 1-benzyloxy(oxalyl), 2-benzyloxy-acetyl, 2-benzenesulfonylamino-ethanoyl, 2-oxo-2-phenyl-ethanoyl, 3*H*-oxazole-2-carbonyl, 5-trifluoromethyl-oxazole-2-carbonyl, 3-trifluoromethyl-[1,2,4]oxadiazole-5-carbonyl, 2,2,3,3,3-pentafluoro-propionyl, hydroxyaminooxalyl, oxalyl, 2-(1,3-dihydro-isindol-2-yl)-2-oxo-acetyl, benzothiazol-2-ylaminooxalyl, 2-oxo-ethyl, 2-oxazol-2-yl-2-oxo-ethyl or 2-benzooxazol-2-yl-2-oxo-ethyl;

X^2 is selected from 5-nitrothiazol-2-ylamino, 2-nitrophenylamino, pyrimidin-2-ylamino, tetrahydro-pyran-4-ylamino, *N*-(2-methoxyethyl)-*N*-(tetrahydro-pyran-4-yl)amino, 1-methyl-piperidin-4-ylamino, isopropylamino, di(thien-2-ylmethyl)amino or di(benzyl)amino; and

R^3 is thiophene-2-sulfonyl-methyl, 3-chloro-2-fluoro-phenyl-methane-sulfonyl-methyl, benzene-sulfonyl-methyl, phenyl-methane-sulfonyl-methyl, 2-(1,1-difluoro-methoxy)-phenyl-methane-sulfonyl-methyl, 2-benzene-sulfonyl-ethyl, 2-(pyridine-2-sulfonyl)-ethyl, 2-(pyridine-4-sulfonyl)-ethyl, 2-phenyl-methanesulfonyl-ethyl, oxy-pyridin-2-yl-methane-sulfonyl-methyl, prop-2-ene-1-sulfonyl-methyl, 4-methoxy-phenyl-methane-sulfonyl-methyl, *p*-tolyl-methane-sulfonyl-methyl, 4-chloro-phenyl-methane-sulfonyl-methyl, *o*-tolyl-methane-sulfonyl-methyl, 3,5-dimethyl-phenyl-methane-sulfonyl-methyl, 4-trifluoromethyl-phenyl-methane-sulfonyl-methyl, 4-trifluoro-methoxy-phenyl-methane-sulfonyl-methyl, 2-bromo-phenyl-methane-sulfonyl-methyl, pyridin-2-yl-methane-sulfonyl-methyl, pyridin-3-yl-methane-sulfonyl-methyl, pyridin-4-yl-methane-sulfonyl-methyl, naphthalen-2-yl-methane-sulfonyl-methyl, 3-methyl-phenyl-methane-sulfonyl-methyl, 3-trifluoro-methyl-phenyl-methane-sulfonyl-methyl, 3-trifluoromethoxy-phenyl-methane-sulfonyl-methyl, 4-fluoro-2-trifluoromethoxy-phenyl-methane-sulfonylmethyl, 2-fluoro-6-trifluoromethyl-phenylmethanesulfonylmethyl, 3-chloro-phenylmethanesulfonylmethyl, 2-fluoro-phenylmethanesulfonylmethyl, 2-trifluoro-phenylmethanesulfonylmethyl, 2-cyano-phenylmethanesulfonylmethyl, 4-*tert*-butyl-phenylmethanesulfonylmethyl, 2-fluoro-3-methyl-phenyl-methane-sulfonyl-methyl, 3-fluoro-phenylmethanesulfonylmethyl, 4-fluoro-phenylmethane-sulfonylmethyl, 2-chloro-phenylmethanesulfonylmethyl, 2,5-difluoro-phenylmethane-

-177-

sulfonylmethyl, 2,6-difluoro-phenylmethanesulfonylmethyl, 2,5-dichloro-phenyl-methane-sulfonylmethyl, 3,4-dichloro-phenylmethanesulfonylmethyl,
2-(1,1-difluoro-methoxy)-phenyl-methanesulfonylmethyl, 2-cyano-phenyl-methane-sulfonyl-methyl, 3-cyano-phenylmethanesulfonylmethyl, 2-trifluoro-methoxy-phenyl-
5 methane-sulfonylmethyl, 2,3-difluoro-phenylmethanesulfonylmethyl,
2,5-difluoro-phenyl-methanesulfonylmethyl, biphenyl-2-ylmethanesulfonylmethyl, cyclohexylmethyl, 3-fluoro-phenyl-methanesulfonylmethyl, 3,4-difluoro-phenyl-methanesulfonylmethyl, 2,4-difluoro-phenylmethanesulfonylmethyl, 2,4,6-trifluoro-phenylmethanesulfonylmethyl, 2,4,5-trifluoro-phenylmethanesulfonylmethyl,
10 2,3,4-trifluoro-phenylmethanesulfonylmethyl, 2,3,5-trifluoro-phenyl-methane-sulfonylmethyl, 2,5,6-trifluoro-phenylmethanesulfonylmethyl, 2-chloro-5-trifluoro-methylphenylmethanesulfonylmethyl, 2-methyl-propane-1-sulfonyl,
2-fluoro-3-trifluoro-methylphenylmethanesulfonylmethyl, 2-fluoro-4-trifluoro-methylphenylmethanesulfonylmethyl, 2-fluoro-5-trifluoro-methyl-phenyl-methane-sulfonyl-methyl, 4-fluoro-3-trifluoro-methylphenylmethanesulfonylmethyl,
15 2-methoxy-phenyl-methanesulfonylmethyl,
3,5-bis-trifluoromethyl-phenylmethanesulfonylmethyl,
4-difluoromethoxy-phenylmethanesulfonylmethyl, 2-difluoro-methoxy-phenyl-methanesulfonylmethyl, 3-difluoromethoxy-phenylmethanesulfonylmethyl,
20 2,6-dichloro-phenylmethanesulfonylmethyl, biphenyl-4-ylmethanesulfonylmethyl, 3,5-dimethyl-isoxazol-4-ylmethanesulfonylmethyl, 5-chloro-thien-2-yl-methane-sulfonylmethyl, 2-[4-(1,1-difluoro-methoxy)-benzenesulfonyl]-ethyl,
2-[2-(1,1-difluoro-methoxy)-benzenesulfonyl]-ethyl, 2-[3-(1,1-difluoro-methoxy)-benzenesulfonyl]-ethyl, 2-(4-trifluoromethoxy-benzenesulfonyl)-ethyl,
25 2-(3-trifluoromethoxy-benzenesulfonyl)-ethyl, 2-(2-trifluoro-methoxy-benzene-sulfonyl)-ethyl, (cyanomethyl-methyl-carbamoyl)-methyl, biphenyl-3-ylmethyl, 2-oxo-2-pyrrolidin-1-yl-ethyl, 2-benzenesulfonyl-ethyl, isobutylsulfanylmethyl, 2-phenylsulfanyl-ethyl, cyclohexylmethanesulfonylmethyl, 2-cyclohexyl-ethanesulfonyl, benzyl, naphthalen-2-yl, benzylsulfanylmethyl,
30 2-trifluoromethyl-benzylsulfanylmethyl, phenylsulfanyl-ethyl, cyclopropyl-methanesulfonylmethyl, 5-bromo-thien-2-ylmethyl, 3-phenyl-propyl, 2,2-difluoro-3-phenyl-propyl, 3,4,5-trimethoxy-phenylmethanesulfonylmethyl, 2,2-difluoro-3-thien-2-yl-propyl, cyclohexylethyl, cyclohexylmethyl, *tert*-butylmethyl,

-178-

1-methylcyclohexylmethyl, 1-methylcyclopentylmethyl, 2,2-difluoro-3-phenylpropyl, 2,2-dimethyl-3-phenylpropyl, 1-benzylcyclopropylmethyl, $-X^5S(O)_2R^{13}$ and $-X^5S(O)_2R^{14}$, wherein R^{13} is alkyl and R^{14} is phenyl which phenyl is unsubstituted or substituted; and the *N*-oxide derivatives, prodrug derivatives, protected derivatives, individual isomers and mixtures of isomers thereof; and the pharmaceutically acceptable salts and solvates of such compounds and the *N*-oxide derivatives, prodrug derivatives, protected derivatives, individual isomers and mixtures of isomers thereof.

13. A compound of Claim 12 in which:

X^3 is 1*H*-benzoimidazol-2-ylcarbonyl, pyrimidin-2-ylcarbonyl, benzooxazol-2-ylcarbonyl, benzothiazol-2-ylcarbonyl, pyridazin-3-ylcarbonyl, 3-phenyl-[1,2,4]oxadiazol-5-ylcarbonyl or 3-ethyl-[1,2,4]oxadiazol-5-ylcarbonyl, 2-oxo-2-pyrrolidin-1-yl-acetyl, 2-morpholin-4-yl-2-oxo-acetyl, 2-oxo-2-piperazin-1-yl-acetyl, 2-(4-methanesulfonyl-piperazin-1-yl)-2-oxo-acetyl, 2-(1,1-dioxo-1*λ*⁶-thiomorpholin-4-yl)-2-oxo-acetyl, dimethylaminooxalyl, tetrahydro-pyran-4-ylaminooxalyl, 2-morpholin-4-yl-ethylaminooxalyl, cyclopentyl-ethylaminooxalyl, pyridin-3-ylaminooxalyl, phenylaminooxalyl or 1-benzoyl-piperidin-4-ylaminooxalyl;

X^2 is selected from -OH, dimethylcarbamoyloxy, morpholin-4-ylcarbonyloxy, piperidin-1-yl-carbonyloxy, pyrrolidin-1-yl-carbonyloxy, pyrimidin-2-ylamino, tetrahydro-pyran-4-ylamino, 1-methyl-piperidin-4-ylamino, *N*-(2-methoxyethyl)-*N*-(tetrahydro-pyran-4-yl)amino, isopropylamino and cyclohexylamino;

R^3 is cyclohexylethyl, cyclohexylmethyl, *tert*-butylmethyl, 1-methylcyclohexylmethyl, 1-methylcyclopentylmethyl, 2,2-difluoro-3-phenylpropyl, 2,2-dimethyl-3-phenylpropyl, 1-benzylcyclopropylmethyl, $-X^5S(O)_2R^{13}$ or $-X^5S(O)_2R^{14}$, wherein R^{13} is alkyl and R^{14} is phenyl which phenyl is unsubstituted or substituted; and the pharmaceutically acceptable salts and solvates of such compounds and the *N*-oxide derivatives, prodrug derivatives, protected derivatives, individual isomers and mixtures of isomers thereof.

14. A compound of Claim 1 selected from the group consisting of:

(*R*)-*N*-cyanomethyl-2-hydroxy-3-phenylmethanesulfonyl-propionamide;

(*R*)-*N*-(1-cyano-1-thiophen-2-yl-methyl)-2-hydroxy-3-phenylmethanesulfonyl-propionamide;

-179-

- (*R*)-*N*-(1-cyano-1-thiophen-2-yl-methyl)-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionamide;
- (*R*)-*N*-cyanomethyl-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionamide;
- morpholine-4-carboxylic acid (*R*)-1-(cyanomethyl-carbamoyl)-2-phenylmethanesulfonyl-ethyl ester;
- 5 morpholine-4-carboxylic acid (*R*)-1-(cyanomethyl-carbamoyl)-2-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-ethyl ester;
- (*R*)-(2-methoxy-ethyl)-carbamic acid 1-(cyanomethyl-carbamoyl)-2-phenylmethanesulfonyl-ethyl ester;
- (*S*)-diethyl-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- 10 (*S*)-pyrrolidine-1-carboxylic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (*S*)-morpholine-4-carboxylic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (*S*)-4-Ethyl-piperazine-1-carboxylic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (*S*)-2-hydroxymethyl-pyrrolidine-1-carboxylic acid (*S*)-1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- 15 (*S*)-(2,2,2-Trifluoro-ethyl)-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (*S*)-(2-hydroxyethyl)-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (Tetrahydrofuran-2-ylmethyl)-carbamic acid (*S*)-1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (*S*)-Azetidine-1-carboxylic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (*S*)-cyclopropyl-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- 20 (*S*)-piperidine-1-carboxylic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (*S*)-(2-methoxy-ethyl)-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (*R*)-3-hydroxy-pyrrolidine-1-carboxylic acid (*S*)-1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (*S*)-3-hydroxy-pyrrolidine-1-carboxylic acid (*S*)-1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (*S*)-morpholine-4-carboxylic acid 1-(cyanomethyl-carbamoyl)-3-cyclohexyl-propyl ester;
- 25 morpholine-4-carboxylic acid (*R*)-1-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester;
- morpholine-4-carboxylic acid (*R*)-1-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propylcarbamoyl]-2-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-ethyl ester;
- morpholine-4-carboxylic acid (*R*)-1-[(*S*)-1-(1-benzothiazol-2-yl-methanoyl)-propylcarbamoyl]-2-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-ethyl ester;
- 30 pyrrolidine-1-carboxylic acid (*R*)-1-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester;
- dimethyl-carbamic acid (*R*)-1-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester;
- 35 morpholine-4-carboxylic acid (*R*)-1-[(*S*)-1-(1-benzylcarbamoyl-methanoyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester;

-180-

- morpholine-4-carboxylic acid (S)-1-[(S)-1-(oxazolo[4,5-b]pyridine-2-carbonyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester;
- morpholine-4-carboxylic acid (S)-1-[(S)-1-(5-ethyl-[1,3,4]oxadiazole-2-carbonyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester;
- 5 (S)-2-[(R)-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propanoylamino]-*N*-methoxy-*N*-methyl-butylamide;
- (R)-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-*N*-[(S)-1-formyl-propyl]-2-hydroxy-propionamide;
- (R)-*N*-[(S)-1-(1-benzooxazol-2-yl-methanoyl)-propyl]-2-hydroxy-3-phenyl-methanesulfonyl-
- 10 propionamide;
- (S)-3-{3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-propanoylamino}-2-oxo-pentanoic acid benzylamide;
- N*-[(S)-1-(1-benzooxazol-2-yl-methanoyl)-propyl]-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-propionamide;
- 15 *N*-[(S)-1-(1-benzooxazol-2-yl-methanoyl)-3-phenyl-propyl]-3-*p*-tolylmethanesulfonyl-propionamide;
- 3-(2-difluoromethoxy-phenylmethanesulfonyl)-*N*-(1-ethyl-2,3-dioxo-3-pyrrolidin-1-yl-propyl)-propionamide;
- 3-(2-difluoromethoxy-phenylmethanesulfonyl)-*N*-(1-ethyl-3-morpholin-4-yl-2,3-dioxo-propyl)-propionamide;
- 20 3-(2-difluoromethoxy-phenylmethanesulfonyl)-*N*-(1-ethyl-2,3-dioxo-3-piperazin-1-yl-propyl)-propionamide;
- 3-(2-difluoromethoxy-phenylmethanesulfonyl)-*N*-[3-(1,1-dioxo-1,6-thiomorpholin-4-yl)-1-ethyl-2,3-dioxo-propyl]-propionamide;
- 3-(2-difluoromethoxy-phenylmethanesulfonyl)-*N*-[1-ethyl-3-(4-methyl-sulfonyl-piperazin-1-yl)-2,3-
- 25 dioxo-propyl]-propionamide;
- 3-[3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid dimethylamide;
- 3-[3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid cyclopentyl-ethyl-amide;
- 30 3-[3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid phenylamide;
- 3-[3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid pyridin-3-ylamide;
- 3-[3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid (tetrahydro-
- 35 pyran-4-yl)-amide;
- 3-[3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid (1-benzoyl-

-181-

- piperidin-4-yl)-amide;
 3-[3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid (2-morpholin-4-yl-ethyl)-amide;
 (R)-N-[(S)-1-(1-benzooxazol-2-yl-methanoyl)-propyl]-2-(2-nitro-phenylamino)-3-phenylmethanesulfonyl-propionamide;
 5 N-[1-(benzooxazole-2-carbonyl)-propyl]-3-phenylmethanesulfonyl-2-(pyrimidin-2-ylamino)-propionamide.
 (R)-N-[(S)-1-(1-benzooxazol-2-yl-methanoyl)-butyl]-2-(5-nitro-thiazol-2-ylamino)-3-phenylmethanesulfonyl-propionamide;
 10 (2S) (4,4-difluoro-2-hydroxy-5-phenyl-pentanoic acid (1(S)-cyano-3-phenyl-propyl)-amide;
 N-(1(S)-cyano-3-phenyl-propyl)-2-(S)-(2-morpholin-4-yl-2-oxo-ethoxy)-4-phenyl-butyramide;
 N-(1-(S)-cyano-3-phenyl-propyl)-2-(S)-fluoro-4-phenyl-butyramide;
 N-(1-(S)-cyano-3-phenyl-propyl)-2,2-difluoro-4-phenyl-butyramide;
 N-(1-(S)-cyano-3-phenyl-propyl)-2-(S)-hydroxy-4-phenyl-butyramide;
 15 N-(1-(S)-cyano-3-phenyl-propyl)-2-(R)-hydroxy-4-phenyl-butyramide;
 N-(1-(S)-cyano-3-phenyl-propyl)-2-(R)-methoxy-4-phenyl-butyramide;
 2,2-difluoro-5-phenyl-pentanoic acid (1-cyano-cyclopropyl)-amide;
 N-(1-(S)-cyano-3-phenyl-propyl)-4-phenyl-butyramide;
 2,2-difluoro-5-phenyl-pentanoic acid ((S)-1-cyano-3-phenyl-propyl)-amide;
 20 N-(4-cyano-1-ethyl-piperidin-4-yl)-3-cyclohexyl-propionamide;
 N-(4-cyano-1-ethyl-piperidin-4-yl)-3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionamide;
 (S)-tert-butyl-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
 (R)-carbamic acid 1-(cyanomethyl-carbamoyl)-2-(2-difluoromethoxy-phenylmethanesulfonyl)-ethyl ester;
 25 (S)-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
 (R)-morpholine-4-carboxylic acid 1-(1-cyano-cyclopropylcarbamoyl)-2-phenylmethanesulfonyl-ethyl ester;
 (R)-morpholine-4-carboxylic acid 1-(4-cyano-tetrahydro-pyran-4-ylcarbamoyl)-2-phenylmethanesulfonyl-ethyl ester;
 30 3-cyclohexyl-2-hydroxy-N-[1-(oxazolo[4,5-b]pyridine-2-carbonyl)-propyl]-propionamide;
 (R)-N-[1-(benzothiazole-2-carbonyl)-butyl]-2-isopropylamino-3-phenylmethanesulfonyl-propionamide;
 (R)-N-[1-(benzothiazole-2-carbonyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
 35 (R)-N-[1-(benzothiazole-2-carbonyl)-butyl]-2-dibenzylamino-3-phenylmethanesulfonyl-propionamide;

-182-

- (R)-N-[1-(benzothiazole-2-carbonyl)-butyl]-2-dimethylamino-3-phenylmethanesulfonyl-propionamide;
- (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- 5 (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-(1-methyl-piperidin-4-ylamino)-3-phenylmethanesulfonyl-propionamide;
- (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-(bis-thiophen-2-ylmethyl-amino)-3-phenylmethanesulfonyl-propionamide;
- (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-dibenzylamino-3-phenylmethanesulfonyl-propionamide;
- 10 (S)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-(tetrahydro-pyran-4-ylamino)-3-thiophen-2-yl-propionamide;
- (S)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-isopropylamino-3-thiophen-2-yl-propionamide;
- (R)-N-[1-(benzothiazole-2-carbonyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- 15 (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-isopropylamino-3-phenylmethanesulfonyl-propionamide;
- 20 (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-[(2-methoxy-ethyl)-(tetrahydro-pyran-4-yl)-amino]-3-phenylmethanesulfonyl-propionamide;
- (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-cyclohexylamino-3-phenylmethanesulfonyl-propionamide;
- (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-dimethylamino-3-phenylmethanesulfonyl-propionamide;
- 25 (1S)-N-[1-(benzoxazole-2-carbonyl)-butyl]-2-(S)-fluoro-4-phenyl-butyramide;
- 2,2-difluoro-5-phenyl-pentanoic acid [(S)-1-(benzoxazole-2-carbonyl)-butyl]-amide;
- morpholine-4-carboxylic acid (S)-1-[(S)-1-(benzoxazole-2-carbonyl)-propylcarbamoyl]-2-cyclohexyl-ethyl ester;
- 30 morpholine-4-carboxylic acid (S)-2-cyclohexyl-1-[(S)-1-(oxazolo[4,5-b]pyridine-2-carbonyl)-propylcarbamoyl]-ethyl ester;
- morpholine-4-carboxylic acid (S)-2-cyclohexyl-1-[(S)-1-(5-ethyl-[1,3,4]oxadiazole-2-carbonyl)-propylcarbamoyl]-ethyl ester;
- morpholine-4-carboxylic acid (S)-2-cyclohexyl-1-[(S)-1-(5-phenyl-[1,3,4]oxadiazole-2-carbonyl)-propylcarbamoyl]-ethyl ester;
- 35

-183-

- morpholine-4-carboxylic acid (S)-1-[(S)-1-(benzoxazole-2-carbonyl)-propylcarbamoyl]-3-cyclohexyl-propyl ester;
- 4-[4,4-dimethyl-2-(morpholine-4-carbonyloxy)-pentanoylamino]-3-oxo-azepane-1-carboxylic acid benzyl ester;
- 5 (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-3-cyclopropylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- (R)-N-[1-(benzoxazole-2-carbonyl)-butyl]-2-cyclohexylamino-3-cyclopropylmethanesulfonyl-propionamide;
- (R)-N-[1-(benzoxazole-2-carbonyl)-butyl]-2-cycloheptylamino-3-cyclopropylmethanesulfonyl-propionamide;
- 10 (R)-3-phenylmethanesulfonyl-N-[(S)-3-phenyl-1-(thiazole-2-carbonyl)-propyl]-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-3-phenyl-propyl]-3-cyclopropylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- 15 (R)-3-cyclopropylmethanesulfonyl-N-[1-(5-ethyl-1,2,4-oxadiazole-3-carbonyl)-propyl]-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- (R)-3-phenylmethanesulfonyl-N-[1-(3-phenyl-1,2,4-oxadiazole-5-carbonyl)-propyl]-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- (R)-N-[1-(3-cyclopropyl-1,2,4-oxadiazole-5-carbonyl)-propyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- 20 {(R)-1-[1-(benzothiazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- {(R)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- 25 {(S)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-thiophen-2-yl-ethyl}-carbamic acid tert-butyl ester;
- {(R)-1-[1-(benzothiazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- {(R)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- 30 {(R)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-cyclopropylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- (R)-1-{1-[hydroxy-(3-phenyl-1,2,4-oxadiazol-5-yl)-methyl]-propylcarbamoyl}-2-phenylmethanesulfonyl-ethyl)-carbamic acid tert-butyl ester;
- 35 ((R)-2-cyclopropylmethanesulfonyl-1-[(S)-1-[(5-ethyl-1,2,4-oxadiazol-3-yl)-hydroxy-methyl]-propylcarbamoyl]-ethyl)-carbamic acid tert-butyl ester;

-184-

- {{(R)-1-[1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamamic acid tert-butyl ester;
- {{(R)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-3-phenyl-propylcarbamoyl]-2-cyclopropylmethanesulfonyl-ethyl}-carbamamic acid tert-butyl ester;
- 5 {{(R)-1-[(S)-1-(hydroxy-thiazol-2-yl-methyl)-3-phenyl-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamamic acid tert-butyl ester;
- {{(R)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-cyclopropylmethanesulfonyl-ethyl}-carbamamic acid tert-butyl ester;
- (R)-1-{1-[hydroxy-(3-phenyl-1,2,4-oxadiazol-5-yl)-methyl]-propylcarbamoyl}-2-phenylmethanesulfonyl-ethyl)-carbamamic acid tert-butyl ester;
- 10 ((R)-2-cyclopropylmethanesulfonyl-1-[(S)-1-[(5-ethyl-1,2,4-oxadiazol-3-yl)-hydroxy-methyl]-propylcarbamoyl]-ethyl)-carbamamic acid tert-butyl ester;
- {{(R)-1-[1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamamic acid tert-butyl ester;
- 15 {{(R)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-3-phenyl-propylcarbamoyl]-2-cyclopropylmethanesulfonyl-ethyl}-carbamamic acid tert-butyl ester;
- {{(R)-1-[(S)-1-(hydroxy-thiazol-2-yl-methyl)-3-phenyl-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamamic acid tert-butyl ester;
- (R)-2-phenylmethanesulfonyl-1-[(S)-1-[(3-cyclopropyl-1,2,4-oxadiazol-5-yl)-hydroxy-methyl]-propylcarbamoyl]-ethyl)-carbamamic acid tert-butyl ester;
- 20 (R)-N-[1-(Benzoxazole-2-carbonyl)-butyl]-2-[cyclopropylmethyl-(tetrahydro-pyran-4-ylmethyl)-amino]-3-phenylmethanesulfonyl-propionamide;
- (R)-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-2-dibenzylamino-3-phenylmethanesulfonyl-propionamide;
- 25 (R)-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- (R)-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-2-isopropylamino-3-phenylmethanesulfonyl-propionamide;
- (R)-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-2-dimethylamino-3-phenylmethanesulfonyl-propionamide;
- 30 (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-(1-methyl-piperidin-4-ylamino)-3-phenylmethanesulfonyl-propionamide;
- 35 (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-(bis-thiophen-2-ylmethyl-amino)-3-phenylmethanesulfonyl-propionamide;

-185-

- (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-dibenzylamino-3-phenylmethanesulfonyl-propionamide;
- (S)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-(tetrahydro-pyran-4-ylamino)-3-thiophen-2-yl-propionamide;
- 5 S)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-isopropylamino-3-thiophen-2-yl-propionamide;
- (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-isopropylamino-3-phenylmethanesulfonyl-propionamide;
- (R)-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- 10 R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- 15 (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-[(2-methoxy-ethyl)-(tetrahydro-pyran-4-yl)-amino]-3-phenylmethanesulfonyl-propionamide;
- (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-cyclohexylamino-3-phenylmethanesulfonyl-propionamide;
- (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-dimethylamino-3-phenylmethanesulfonyl-propionamide;
- 20 N-cyanomethyl-3-cyclohexyl-propionamide;
- N-cyanomethyl-3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionamide;
- 3-(3-cyclohexyl-propionylamino)-2-oxo-5-phenyl-pentanoic acid thiazol-2-ylamide;
- 3-cyclohexyl-N-(1-formyl-3-phenyl-propyl)-propionamide;
- 25 3-(2-difluoromethoxy-phenylmethanesulfonyl)-N-[(S)-1-(5-ethyl-[1,3,4]oxadiazole-2-carbonyl)-propyl]-propionamide;
- N-[(S)-1-(benzoxazol-2-carbonyl)-propyl]-2-(2-cyano-phenylamino)-3-cyclohexyl-propionamide;
- N-Cyanomethyl-3-cyclohexyl-2-(4-methoxy-phenoxy)-propionamide;
- 2-benzyloxy-N-cyanomethyl-3-cyclohexyl-propionamide;
- 30 (R)-N-[(S)-1-(1-benzoxazol-2-yl-methanoyl)-butyl]-2-benzyloxy-3-phenylmethanesulfonyl-propionamide;
- (R)-N-[(S)-1-(1-benzoxazol-2-yl-methanoyl)-propyl]-2-methoxymethoxy-3-phenylmethanesulfonyl-propionamide;
- (S)-N-[(S)-1-(1-benzoxazol-2-yl-methanoyl)-butyl]-2-hydroxy-3-phenyl-propionamide;
- 35 (R)-N-[(S)-1-(1-benzoxazol-2-yl-methanoyl)-propyl]-3-phenylmethanesulfonyl-2-triisopropylsilyloxy-propionamide;

-186-

- (*R*)-*N*-[(*S*)-1-(1-benzothiazol-2-yl-methanoyl)-propyl]-2-hydroxy-3-phenylmethanesulfonyl-propionamide;
- (*R*)-2-hydroxy-3-phenylmethanesulfonyl-*N*-[(*S*)-1-(1-pyridazin-3-yl-methanoyl)-butyl]-propionamide;
- (*S*)-3-[(*R*)-2-hydroxy-3-phenylmethanesulfonyl-propanoylamino]-2-oxo-pentanoic acid benzylamide;
- 5 (*R*)-*N*-[(*S*)-1-(1-benzoxazol-2-yl-methanoyl)-propyl]-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionamide;
- (*R*)-*N*-[(*S*)-1-(1-benzothiazol-2-yl-methanoyl)-propyl]-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionamide;
- (2*R*,5*S*)-2-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonylmethyl]-6-ethoxy-5-ethyl-morpholin-3-one; and their corresponding N-oxides, and their prodrugs, and their protected derivatives, individual
- 10 isomers and mixtures of isomers thereof; and the pharmaceutically acceptable salts and solvates (e.g. hydrates) of such compounds and their N-oxides and their prodrugs, and their protected derivatives, individual isomers and mixtures of isomers thereof.
- 15 15. A compound of claim 14 selected from the group consisting of:
- (*R*)-*N*-cyanomethyl-2-hydroxy-3-phenylmethanesulfonyl-propionamide;
- (*R*)-*N*-(1-cyano-1-thiophen-2-yl-methyl)-2-hydroxy-3-phenylmethanesulfonyl-propionamide;
- (*R*)-*N*-(1-cyano-1-thiophen-2-yl-methyl)-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionamide;
- 20 (*R*)-*N*-cyanomethyl-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionamide;
- morpholine-4-carboxylic acid (*R*)-1-(cyanomethyl-carbamoyl)-2-phenylmethanesulfonyl-ethyl ester;
- morpholine-4-carboxylic acid (*R*)-1-(cyanomethyl-carbamoyl)-2-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-ethyl ester;
- (*R*)-(2-methoxy-ethyl)-carbamic acid 1-(cyanomethyl-carbamoyl)-2-phenylmethanesulfonyl-ethyl
- 25 ester;
- (*S*)-diethyl-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (*S*)-pyrrolidine-1-carboxylic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (*S*)-morpholine-4-carboxylic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (*S*)-4-Ethyl-piperazine-1-carboxylic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- 30 (*S*)-2-hydroxymethyl-pyrrolidine-1-carboxylic acid (*S*)-1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (*S*)-(2,2,2-Trifluoro-ethyl)-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (*S*)-(2-hydroxyethyl)-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (Tetrahydrofuran-2-ylmethyl)-carbamic acid (*S*)-1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- 35 (*S*)-Azetidine-1-carboxylic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
- (*S*)-cyclopropyl-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;

- (S)-piperidine-1-carboxylic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
 (S)-(2-methoxy-ethyl)-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
 (R)-3-hydroxy-pyrrolidine-1-carboxylic acid (S)-1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
 (S)-3-hydroxy-pyrrolidine-1-carboxylic acid (S)-1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
 5 (S)-morpholine-4-carboxylic acid 1-(cyanomethyl-carbamoyl)-3-cyclohexyl-propyl ester;
 morpholine-4-carboxylic acid (*R*)-1-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester;
 morpholine-4-carboxylic acid (*R*)-1-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propylcarbamoyl]-2-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-ethyl ester;
 10 morpholine-4-carboxylic acid (*R*)-1-[(*S*)-1-(1-benzothiazol-2-yl-methanoyl)-propylcarbamoyl]-2-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-ethyl ester;
 pyrrolidine-1-carboxylic acid (*R*)-1-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester;
 dimethyl-carbamic acid (*R*)-1-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester;
 15 morpholine-4-carboxylic acid (*R*)-1-[(*S*)-1-(1-benzylcarbamoyl-methanoyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester;
 morpholine-4-carboxylic acid (S)-1-[(*S*)-1-(oxazolo[4,5-*b*]pyridine-2-carbonyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester;
 20 morpholine-4-carboxylic acid (S)-1-[(*S*)-1-(5-ethyl-[1,3,4]oxadiazole-2-carbonyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester;
 (S)-2-[(*R*)-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propanoylamino]-*N*-methoxy-*N*-methyl-butylamide;
 (R)-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-*N*-[(*S*)-1-formyl-propyl]-2-hydroxy-propionamide;
 25 (R)-*N*-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propyl]-2-hydroxy-3-phenyl-methanesulfonyl-propionamide;
 (S)-3-{3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-propanoylamino}-2-oxo-pentanoic acid benzylamide;
 30 *N*-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propyl]-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-propionamide;
N-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-3-phenyl-propyl]-3-*p*-tolylmethanesulfonyl-propionamide;
 3-(2-difluoromethoxy-phenylmethanesulfonyl)-*N*-(1-ethyl-2,3-dioxo-3-pyrrolidin-1-yl-propyl)-propionamide;
 35 3-(2-difluoromethoxy-phenylmethanesulfonyl)-*N*-(1-ethyl-3-morpholin-4-yl-2,3-dioxo-propyl)-propionamide;

-188-

- 3-(2-difluoromethoxy-phenylmethanesulfonyl)-*N*-(1-ethyl-2,3-dioxo-3-piperazin-1-yl-propyl)-propionamide;
- 3-(2-difluoromethoxy-phenylmethanesulfonyl)-*N*-[3-(1,1-dioxo-116-thiomorpholin-4-yl)-1-ethyl-2,3-dioxo-propyl]-propionamide;
- 5 3-(2-difluoromethoxy-phenylmethanesulfonyl)-*N*-[1-ethyl-3-(4-methyl-sulfonyl-piperazin-1-yl)-2,3-dioxo-propyl]-propionamide;
- 3-[3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid dimethylamide;
- 3-[3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid cyclopentyl-ethyl-amide;
- 10 3-[3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid phenylamide;
- 3-[3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid pyridin-3-ylamide;
- 15 3-[3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid (tetrahydro-pyran-4-yl)-amide;
- 3-[3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid (1-benzoyl-piperidin-4-yl)-amide;
- 3-[3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionylamino]-2-oxo-pentanoic acid (2-morpholin-4-yl-ethyl)-amide;
- 20 (R)-*N*-[(S)-1-(1-benzooxazol-2-yl-methanoyl)-propyl]-2-(2-nitro-phenylamino)-3-phenylmethanesulfonyl-propionamide;
- N*-[1-(benzooxazole-2-carbonyl)-propyl]-3-phenylmethanesulfonyl-2-(pyrimidin-2-ylamino)-propionamide.
- 25 (R)-*N*-[(S)-1-(1-benzooxazol-2-yl-methanoyl)-butyl]-2-(5-nitro-thiazol-2-ylamino)-3-phenylmethanesulfonyl-propionamide;
- (2S) (4,4-difluoro-2-hydroxy-5-phenyl-pentanoic acid (1(S)-cyano-3-phenyl-propyl)-amide;
- N*-(1(S)-cyano-3-phenyl-propyl)-2-(S)-(2-morpholin-4-yl-2-oxo-ethoxy)-4-phenyl-butyramide;
- N*-(1(S)-cyano-3-phenyl-propyl)-2-(S)-fluoro-4-phenyl-butyramide;
- 30 *N*-(1(S)-cyano-3-phenyl-propyl)-2,2-difluoro-4-phenyl-butyramide;
- N*-(1(S)-cyano-3-phenyl-propyl)-2-(S)-hydroxy-4-phenyl-butyramide;
- N*-(1(S)-cyano-3-phenyl-propyl)-2-(R)-hydroxy-4-phenyl-butyramide;
- N*-(1(S)-cyano-3-phenyl-propyl)-2-(R)-methoxy-4-phenyl-butyramide;
- 2,2-difluoro-5-phenyl-pentanoic acid (1-cyano-cyclopropyl)-amide;
- 35 *N*-(1(S)-cyano-3-phenyl-propyl)-4-phenyl-butyramide;

-189-

- 2,2-difluoro-5-phenyl-pentanoic acid ((S)-1-cyano-3-phenyl-propyl)-amide;
N-(4-cyano-1-ethyl-piperidin-4-yl)-3-cyclohexyl-propionamide;
N-(4-cyano-1-ethyl-piperidin-4-yl)-3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionamide;
(S)-tert-butyl-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
5 (R)-carbamic acid 1-(cyanomethyl-carbamoyl)-2-(2-difluoromethoxy-phenylmethanesulfonyl)-ethyl ester;
(S)-carbamic acid 1-(cyanomethyl-carbamoyl)-2-cyclohexyl-ethyl ester;
(R)-morpholine-4-carboxylic acid 1-(1-cyano-cyclopropylcarbamoyl)-2-phenylmethanesulfonyl-ethyl ester;
10 (R)-morpholine-4-carboxylic acid 1-(4-cyano-tetrahydro-pyran-4-ylcarbamoyl)-2-phenylmethanesulfonyl-ethyl ester;
3-cyclohexyl-2-hydroxy-*N*-[1-(oxazolo[4,5-*b*]pyridine-2-carbonyl)-propyl]-propionamide;
(R)-*N*-[1-(benzothiazole-2-carbonyl)-butyl]-2-isopropylamino-3-phenylmethanesulfonyl-propionamide;
15 (R)-*N*-[1-(benzothiazole-2-carbonyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
(R)-*N*-[1-(benzothiazole-2-carbonyl)-butyl]-2-dibenzylamino-3-phenylmethanesulfonyl-propionamide;
(R)-*N*-[1-(benzothiazole-2-carbonyl)-butyl]-2-dimethylamino-3-phenylmethanesulfonyl-propionamide;
20 (R)-*N*-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
(R)-*N*-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-(1-methyl-piperidin-4-ylamino)-3-phenylmethanesulfonyl-propionamide;
(R)-*N*-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-(bis-thiophen-2-ylmethyl-amino)-3-phenylmethanesulfonyl-propionamide;
25 (R)-*N*-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-dibenzylamino-3-phenylmethanesulfonyl-propionamide;
(S)-*N*-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-(tetrahydro-pyran-4-ylamino)-3-thiophen-2-yl-propionamide;
30 (S)-*N*-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-isopropylamino-3-thiophen-2-yl-propionamide;
(R)-*N*-[1-(benzothiazole-2-carbonyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
(R)-*N*-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
35 (R)-*N*-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-isopropylamino-3-phenylmethanesulfonyl-propionamide;

-190-

- (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-[(2-methoxy-ethyl)-(tetrahydro-pyran-4-yl)-amino]-3-phenylmethanesulfonyl-propionamide;
- (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-cyclohexylamino-3-phenylmethanesulfonyl-propionamide;
- 5 (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-2-dimethylamino-3-phenylmethanesulfonyl-propionamide;
- (1S)-N-[1-(benzoxazole-2-carbonyl)-butyl]-2-(S)-fluoro-4-phenyl-butyramide;
- 2,2-difluoro-5-phenyl-pentanoic acid [(S)-1-(benzoxazole-2-carbonyl)-butyl]-amide;
- morpholine-4-carboxylic acid (S)-1-[(S)-1-(benzoxazole-2-carbonyl)-propylcarbamoyl]-2-
- 10 cyclohexyl-ethyl ester;
- morpholine-4-carboxylic acid (S)-2-cyclohexyl-1-[(S)-1-(oxazolo[4,5-b]pyridine-2-carbonyl)-propylcarbamoyl]-ethyl ester;
- morpholine-4-carboxylic acid (S)-2-cyclohexyl-1-[(S)-1-(5-ethyl-[1,3,4]oxadiazole-2-carbonyl)-propylcarbamoyl]-ethyl ester;
- 15 morpholine-4-carboxylic acid (S)-2-cyclohexyl-1-[(S)-1-(5-phenyl-[1,3,4]oxadiazole-2-carbonyl)-propylcarbamoyl]-ethyl ester;
- morpholine-4-carboxylic acid (S)-1-[(S)-1-(benzoxazole-2-carbonyl)-propylcarbamoyl]-3-cyclohexyl-propyl ester;
- 4-[4,4-dimethyl-2-(morpholine-4-carboxyloxy)-pentanoylamino]-3-oxo-azepane-1-carboxylic acid
- 20 benzyl ester;
- (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-butyl]-3-cyclopropylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- (R)-N-[1-(benzoxazole-2-carbonyl)-butyl]-2-cyclohexylamino-3-cyclopropylmethanesulfonyl-propionamide;
- 25 (R)-N-[1-(benzoxazole-2-carbonyl)-butyl]-2-cycloheptylamino-3-cyclopropylmethanesulfonyl-propionamide;
- (R)-3-phenylmethanesulfonyl-N-[(S)-3-phenyl-1-(thiazole-2-carbonyl)-propyl]-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- (R)-N-[(S)-1-(benzoxazole-2-carbonyl)-3-phenyl-propyl]-3-cyclopropylmethanesulfonyl-2-
- 30 (tetrahydro-pyran-4-ylamino)-propionamide;
- (R)-3-cyclopropylmethanesulfonyl-N-[1-(5-ethyl-1,2,4-oxadiazole-3-carbonyl)-propyl]-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- (R)-3-phenylmethanesulfonyl-N-[1-(3-phenyl-1,2,4-oxadiazole-5-carbonyl)-propyl]-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- 35 (R)-N-[1-(3-cyclopropyl-1,2,4-oxadiazole-5-carbonyl)-propyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;

- {(R)-1-[1-(benzothiazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- {(R)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- 5 {(S)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-thiophen-2-yl-ethyl}-carbamic acid tert-butyl ester;
- {(R)-1-[1-(benzothiazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- {(R)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- 10 {(R)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-cyclopropylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- (R)-1-[1-[hydroxy-(3-phenyl-1,2,4-oxadiazol-5-yl)-methyl]-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl)-carbamic acid tert-butyl ester;
- 15 ((R)-2-cyclopropylmethanesulfonyl-1-[(S)-1-[(5-ethyl-1,2,4-oxadiazol-3-yl)-hydroxy-methyl]-propylcarbamoyl]-ethyl)-carbamic acid tert-butyl ester;
- {(R)-1-[1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- {(R)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-3-phenyl-propylcarbamoyl]-2-cyclopropylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- 20 {(R)-1-[(S)-1-(hydroxy-thiazol-2-yl-methyl)-3-phenyl-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- {(R)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-cyclopropylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- 25 (R)-1-[1-[hydroxy-(3-phenyl-1,2,4-oxadiazol-5-yl)-methyl]-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl)-carbamic acid tert-butyl ester;
- ((R)-2-cyclopropylmethanesulfonyl-1-[(S)-1-[(5-ethyl-1,2,4-oxadiazol-3-yl)-hydroxy-methyl]-propylcarbamoyl]-ethyl)-carbamic acid tert-butyl ester;
- {(R)-1-[1-(benzoxazol-2-yl-hydroxy-methyl)-butylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- 30 {(R)-1-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-3-phenyl-propylcarbamoyl]-2-cyclopropylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- {(R)-1-[(S)-1-(hydroxy-thiazol-2-yl-methyl)-3-phenyl-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl}-carbamic acid tert-butyl ester;
- 35 (R)-2-phenylmethanesulfonyl-1-[(S)-1-[(3-cyclopropyl-1,2,4-oxadiazol-5-yl)-hydroxy-methyl]-propylcarbamoyl]-ethyl)-carbamic acid tert-butyl ester;

-192-

- (R)-N-[1-(Benzoxazole-2-carbonyl)-butyl]-2-[cyclopropylmethyl-(tetrahydro-pyran-4-ylmethyl)-amino]-3-phenylmethanesulfonyl-propionamide;
- (R)-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-2-dibenzylamino-3-phenylmethanesulfonyl-propionamide;
- 5 (R)-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- (R)-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-2-isopropylamino-3-phenylmethanesulfonyl-propionamide;
- (R)-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-2-dimethylamino-3-phenylmethanesulfonyl-propionamide;
- 10 (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-(1-methyl-piperidin-4-ylamino)-3-phenylmethanesulfonyl-propionamide;
- 15 (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-(bis-thiophen-2-ylmethyl-amino)-3-phenylmethanesulfonyl-propionamide;
- (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-dibenzylamino-3-phenylmethanesulfonyl-propionamide;
- (S)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-(tetrahydro-pyran-4-ylamino)-3-thiophen-2-yl-propionamide;
- 20 S)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-isopropylamino-3-thiophen-2-yl-propionamide;
- (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-isopropylamino-3-phenylmethanesulfonyl-propionamide;
- 25 (R)-N-[1-(benzothiazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- 30 (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-[(2-methoxy-ethyl)-(tetrahydro-pyran-4-yl)-amino]-3-phenylmethanesulfonyl-propionamide;
- (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-cyclohexylamino-3-phenylmethanesulfonyl-propionamide;
- 35 (R)-N-[(S)-1-(benzoxazol-2-yl-hydroxy-methyl)-butyl]-2-dimethylamino-3-phenylmethanesulfonyl-propionamide;

-193-

- N*-cyanomethyl-3-cyclohexyl-propionamide;
N-cyanomethyl-3-(2-difluoromethoxy-phenylmethanesulfonyl)-propionamide;
 3-(3-cyclohexyl-propionylamino)-2-oxo-5-phenyl-pentanoic acid thiazol-2-ylamide;
 3-cyclohexyl-*N*-(1-formyl-3-phenyl-propyl)-propionamide;
 5 3-(2-difluoromethoxy-phenylmethanesulfonyl)-*N*-[(*S*)-1-(5-ethyl-[1,3,4]oxadiazole-2-carbonyl)-propyl]-propionamide;
N-[(*S*)-1-(benzooxazole-2-carbonyl)-propyl]-2-(2-cyano-phenylamino)-3-cyclohexyl-propionamide;
N-Cyanomethyl-3-cyclohexyl-2-(4-methoxy-phenoxy)-propionamide;
 2-benzyloxy-*N*-cyanomethyl-3-cyclohexyl-propionamide;
 10 (*R*)-*N*-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-butyl]-2-benzyloxy-3-phenylmethanesulfonyl-propionamide;
 (*R*)-*N*-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propyl]-2-methoxymethoxy-3-phenylmethanesulfonyl-propionamide;
 (*S*)-*N*-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-butyl]-2-hydroxy-3-phenyl-propionamide;
 15 (*R*)-*N*-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propyl]-3-phenylmethanesulfonyl-2-triisopropylsilyloxy-propionamide;
 (*R*)-*N*-[(*S*)-1-(1-benzothiazol-2-yl-methanoyl)-propyl]-2-hydroxy-3-phenylmethanesulfonyl-propionamide;
 (*R*)-2-hydroxy-3-phenylmethanesulfonyl-*N*-[(*S*)-1-(1-pyridazin-3-yl-methanoyl)-butyl]-propionamide;
 20 (*S*)-3-((*R*)-2-hydroxy-3-phenylmethanesulfonyl-propanoylamino)-2-oxo-pentanoic acid benzylamide;
 (*R*)-*N*-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propyl]-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionamide;
 (*R*)-*N*-[(*S*)-1-(1-benzothiazol-2-yl-methanoyl)-propyl]-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionamide; and
 25 (2*R*,5*S*)-2-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonylmethyl]-6-ethoxy-5-ethyl-morpholin-3-one.

16. A compound of claim 15 selected from the group consisting of:

- 30 morpholine-4-carboxylic acid (*R*)-1-(cyanomethyl-carbamoyl)-2-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-ethyl ester, (Compound 31);
 morpholine-4-carboxylic acid (*R*)-1-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester, (Compound 11);
 35 morpholine-4-carboxylic acid (*R*)-1-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propylcarbamoyl]-2-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-ethyl ester, (Compound 14);
 morpholine-4-carboxylic acid (*R*)-1-[(*S*)-1-(1-benzothiazol-2-yl-methanoyl)-propylcarbamoyl]-2-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-ethyl ester, (Compound 15);
 40

-194-

- pyrrolidine-1-carboxylic acid (*R*)-1-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester, (Compound 19);
- 5 dimethyl-carbamic acid (*R*)-1-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester, (Compound 20);
- morpholine-4-carboxylic acid (*R*)-1-[(*S*)-1-(1-benzylcarbamoyl-methanoyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester, (Compound 25);
- 10 morpholine-4-carboxylic acid (*S*)-1-[(*S*)-1-(oxazolo[4,5-*b*]pyridine-2-carbonyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester;
- morpholine-4-carboxylic acid (*S*)-1-[(*S*)-1-(5-ethyl-[1,3,4]oxadiazole-2-carbonyl)-propylcarbamoyl]-2-phenylmethanesulfonyl-ethyl ester;
- 15 (*R*)-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-*N*-[(*S*)-1-formyl-propyl]-2-hydroxy-propionamide;
- (*R*)-*N*-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propyl]-2-hydroxy-3-phenyl-methanesulfonyl-propionamide;
- 20 (*S*)-3-{3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-propanoylamino}-2-oxo-pentanoic acid benzylamide;
- (*R*)-*N*-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propyl]-2-(2-nitro-phenylamino)-3-phenylmethanesulfonyl-propionamide;
- 25 (*R*)-*N*-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-butyl]-2-(5-nitro-thiazol-2-ylamino)-3-phenylmethanesulfonyl-propionamide;
- 30 (*R*)-*N*-[(*S*)-1-(benzoxazole-2-carbonyl)-butyl]-3-phenylmethanesulfonyl-2-(tetrahydro-pyran-4-ylamino)-propionamide;
- (*R*)-*N*-[(*S*)-1-(benzoxazole-2-carbonyl)-butyl]-2-isopropylamino-3-phenylmethanesulfonyl propionamide;
- 35 (*R*)-*N*-[(*S*)-1-(benzoxazole-2-carbonyl)-butyl]-2-[(2-methoxy-ethyl)-(tetrahydro-pyran-4-yl)-amino]-3-phenylmethanesulfonyl-propionamide;
- 40 (*R*)-*N*-[(*S*)-1-(benzoxazole-2-carbonyl)-butyl]-2-cyclohexylamino-3-phenylmethanesulfonyl-propionamide;
- morpholine-4-carboxylic acid (*S*)-2-cyclohexyl-1-[(*S*)-1-(oxazolo[4,5-*b*]pyridine-2-carbonyl)-propylcarbamoyl]-ethyl ester;
- (*S*)-3-[(*R*)-2-hydroxy-3-phenylmethanesulfonyl-propanoylamino]-2-oxo-pentanoic acid benzylamide;
- 45 (*R*)-*N*-[(*S*)-1-(1-benzooxazol-2-yl-methanoyl)-propyl]-3-[2-(1,1-difluoro-methoxy)-phenylmethanesulfonyl]-2-hydroxy-propionamide.

17. A pharmaceutical composition comprising a therapeutically effective amount

-195-

of a compound of Claim 1 in combination with a pharmaceutically acceptable excipient.

5 18. A pharmaceutical composition comprising a therapeutically effective amount of a compound of Claim 2 in combination with a pharmaceutically acceptable excipient.

10 19. A method for treating a disease in an animal in which inhibition of Cathepsin S can prevent, inhibit or ameliorate the pathology and/or symptomology of the disease, which method comprises administering to the animal a therapeutically effective amount of compound of Claim 1 or Claim 2.

15 20. The use of a compound of Claim 1 or 2 in the manufacture of a medicament for treating a disease in an animal in which Cathepsin S activity contributes to the pathology and/or symptomology of the disease.

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
12 December 2002 (12.12.2002)

PCT

(10) International Publication Number
WO 02/098850 A3

(51) International Patent Classification⁷: **C07C 317/46**,
A61K 31/16, A61P 19/02, 33/06, C07C 233/31, 255/29,
255/44, 255/46, 271/12, 271/24, 317/44, C07D 205/04,
207/08, 211/66, 239/08, 263/56, 271/10, C07C 271/16,
C07D 223/08, 277/28, 277/64, 277/82, 295/18, 295/20,
333/24, 307/14, 309/14, 413/12, 413/14, 417/12, C07C
317/46, 255/29, 317/44, A61K 31/16, C07C 255/44,
C07D 205/04, A61P 19/02, C07C 255/44, C07D 207/08,
A61P 33/06, C07C 271/12, C07D 211/66, C07C 233/31,
271/24, C07D 239/08

(21) International Application Number: PCT/US02/17411

(22) International Filing Date: 3 June 2002 (03.06.2002)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
60/295,301 1 June 2001 (01.06.2001) US

(71) Applicants (for all designated States except US): **AXYS
PHARMACEUTICALS, INC.** [US/US]; 180 Kimball
Way, South San Francisco, CA 94080 (US). **AVENTIS
PHARMACEUTICALS INC.** [US/US]; 300 Somerset
Corporate Blvd., Bridgewater, NJ 08807-2854 (US).

(72) Inventors; and

(75) Inventors/Applicants (for US only): **GRAUPE, Michael**
[AT/US]; 1131 Banyan Way, Pacifica, CA 94044 (US).
LI, Jiayao [CN/US]; 888 Foster City Blvd., Foster City,
CA 94404 (US). **LINK, John, O.** [US/US]; 12 Prospect
Ave., San Francisco, CA 94110 (US). **ZIPFEL, Sheila**
[US/US]; Ayy's Pharmaceuticals Inc., 180 Kimball Way,
South San Francisco, CA 94080 (US). **TIMM, Andreas,**
P. [DE/US]; 97 Bonney Court, Bridgewater, NJ 08807
(US). **ALDOUS, David, J.** [US/US]; 31 Dunbridge Lane,

Gillette, NJ 07933 (US). **THURAIRATNAM, Sukan-
thini** [GB/US]; 6 Smoke Rise Lane, Bedminster, NJ 07921
(US).

(74) Agents: **WANG, George, G.** et al.; Aventis Pharmaceu-
ticals Inc., Route 202-206, P.O. Box 6800, Bridgewater, NJ
08807-0800 (US).

(81) Designated States (*national*): AE, AG, AL, AM, AT, AU,
AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU,
CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW,
MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG,
SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ,
VN, YU, ZA, ZM, ZW.

(84) Designated States (*regional*): ARIPO patent (GH, GM,
KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW),
Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM),
European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR,
GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent
(BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR,
NE, SN, TD, TG).

Declaration under Rule 4.17:

— of inventorship (Rule 4.17(iv)) for US only

Published:

— with international search report

(88) Date of publication of the international search report:
24 April 2003

For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
ning of each regular issue of the PCT Gazette.

WO 02/098850 A3

(54) Title: CHEMICAL COMPOUNDS AND PHARMACEUTICAL COMPOSITIONS AS CATHEPSIN S INHIBITORS

(57) Abstract: The present invention relates to compounds of formula (I) (in which X¹ is-NHC(R¹)(R²)X³ or -NHX⁴ and the other variables are as defined in the claims) and the pharmaceutically acceptable salts and N-oxides thereof, useful as selective cathepsin S inhibitors, their uses as therapeutic agents and the methods for their making. Formula (I):

INTERNATIONAL SEARCH REPORT

Int. Application No.

PCT/US 02/17411

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07C317/46 A61K31/16 A61P19/02 A61P33/06 C07C233/31
 C07C255/29 C07C255/44 C07C255/46 C07C271/12 C07C271/24
 C07C317/44 C07D205/04 C07D207/08 C07D211/66 C07D239/08

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07C C07D C07F A61K A61P

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, CHEM ABS Data, BEILSTEIN Data, WPI Data, PAJ

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 00 55144 A (AXYS PHARMACEUTICALS) 21 September 2000 (2000-09-21) page 2 -page 7	1-20
A	WO 01 19796 A (AXYS PHARMACEUTICALS) 22 March 2001 (2001-03-22) page 2 -page 4	1-20
A	WO 01 19808 A (AXYS PHARMACEUTICALS) 22 March 2001 (2001-03-22) page 2 -page 4	1-20

☐ Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

* Special categories of cited documents :

A document defining the general state of the art which is not considered to be of particular relevance

E earlier document but published on or after the international filing date

L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

O document referring to an oral disclosure, use, exhibition or other means

P document published prior to the international filing date but later than the priority date claimed

T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

& document member of the same patent family

Date of the actual completion of the international search

14 January 2003

Date of mailing of the international search report

30/01/2003

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
 NL - 2280 HV Rijswijk
 Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
 Fax: (+31-70) 340-3016

Authorized officer

English, R

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 02/17411

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07D263/56 C07D271/10 C07C271/16 C07D223/08 C07D277/28
C07D277/64 C07D277/82 C07D295/18 C07D295/20 C07D333/24
C07D307/14 C07D309/14 C07D413/12 C07D413/14 C07D417/12

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.

☐ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents :

A document defining the general state of the art which is not considered to be of particular relevance

E earlier document but published on or after the international filing date

L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

O document referring to an oral disclosure, use, exhibition or other means

P document published prior to the international filing date but later than the priority date claimed

T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

Z document member of the same patent family

Date of the actual completion of the international search

14 January 2003

Date of mailing of the international search report

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

English, R

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 02/17411

A. CLASSIFICATION OF SUBJECT MATTER
 IPC 7 C07D498/04 C07F7/18

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.

☐ Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

* Special categories of cited documents:

A document defining the general state of the art which is not considered to be of particular relevance

E earlier document but published on or after the international filing date

L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

O document referring to an oral disclosure, use, exhibition or other means

P document published prior to the international filing date but later than the priority date claimed

T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

& document member of the same patent family

Date of the actual completion of the international search

14 January 2003

Date of mailing of the international search report

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
 NL - 2280 HV Rijswijk
 Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
 Fax: (+31-70) 340-3016

Authorized officer

English, R

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.2

Claims Nos.: 1-15,17,18 (in part)

Present claims 1-13, 17 and 18 relate to an extremely large number of possible compounds or pharmaceutical compositions. Support within the meaning of Article 6 PCT and/or disclosure within the meaning of Article 5 PCT is to be found, however, for only a very small proportion of the compounds or pharmaceutical compositions claimed. In the present case, the claims so lack support, and the application so lacks disclosure, that a meaningful search over the whole of the claimed scope is impossible. Consequently, the search has been carried out for those parts of the claims which appear to be supported and disclosed, namely those parts relating to the compounds prepared in the examples.

The scope of the claims 1-14, in as far as the expressions "prodrug" and "protected derivatives" are concerned, is so unclear (Article 6 PCT) that a meaningful International Search is impossible with regard to these expressions.

The applicant's attention is drawn to the fact that claims, or parts of claims, relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US 02/17411

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

Although claim 19 is directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.
2. ☒ Claims Nos.: 1-15,17,18 (in part)
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:

see FURTHER INFORMATION sheet PCT/ISA/210
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Int. Application No

PCT/US 02/17411

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
WO 0055144	A	21-09-2000	AU 3750700 A	04-10-2000
			BG 105969 A	31-05-2002
			BR 0009044 A	15-01-2002
			CN 1345314 T	17-04-2002
			CZ 20013247 A3	15-05-2002
			EP 1161422 A1	12-12-2001
			HU 0200572 A2	29-06-2002
			NO 20014483 A	01-11-2001
			SK 12872001 A3	09-05-2002
			TR 200103335 T2	22-04-2002
			WO 0055144 A1	21-09-2000
WO 0119796	A	22-03-2001	AU 7490900 A	17-04-2001
			AU 7703300 A	17-04-2001
			EP 1212302 A1	12-06-2002
			WO 0119808 A1	22-03-2001
			WO 0119796 A1	22-03-2001
			US 6492362 B1	10-12-2002
WO 0119808	A	22-03-2001	AU 7490900 A	17-04-2001
			AU 7703300 A	17-04-2001
			EP 1212302 A1	12-06-2002
			WO 0119808 A1	22-03-2001
			WO 0119796 A1	22-03-2001
			US 6492362 B1	10-12-2002

**This Page is Inserted by IFW Indexing and Scanning
Operations and is not part of the Official Record**

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images include but are not limited to the items checked:

- ☐ **BLACK BORDERS**
- ☐ **IMAGE CUT OFF AT TOP, BOTTOM OR SIDES**
- ☐ **FADED TEXT OR DRAWING**
- ☐ **BLURRED OR ILLEGIBLE TEXT OR DRAWING**
- ☐ **SKEWED/SLANTED IMAGES**
- ☒ **COLOR OR BLACK AND WHITE PHOTOGRAPHS**
- ☒ **GRAY SCALE DOCUMENTS**
- ☒ **LINES OR MARKS ON ORIGINAL DOCUMENT**
- ☐ **REFERENCE(S) OR EXHIBIT(S) SUBMITTED ARE POOR QUALITY**
- ☐ **OTHER:** _____

IMAGES ARE BEST AVAILABLE COPY.

As rescanning these documents will not correct the image problems checked, please do not report these problems to the IFW Image Problem Mailbox.